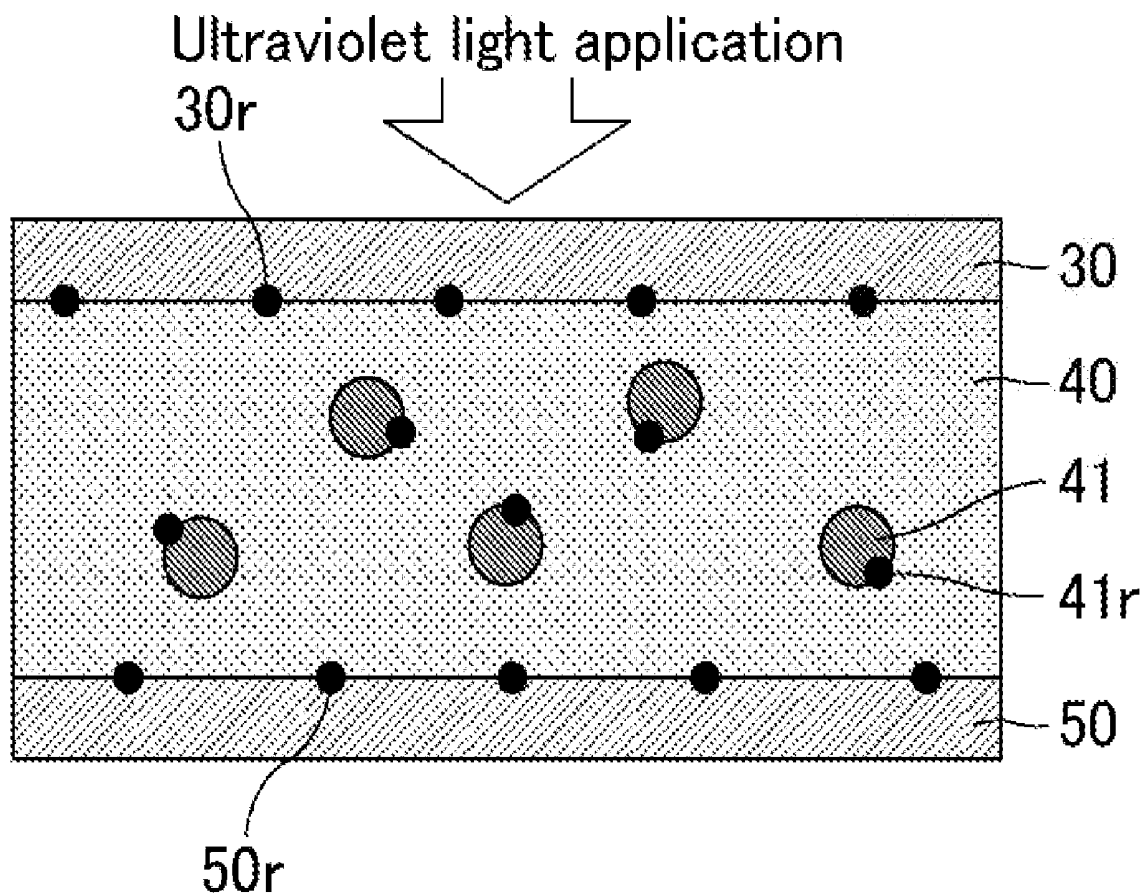




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MIZUSAKI(10) **Pub. No.: US 2020/0026129 A1**(43) **Pub. Date: Jan. 23, 2020**(54) **LIQUID CRYSTAL DISPLAY DEVICE AND
METHOD FOR MANUFACTURING LIQUID
CRYSTAL DISPLAY DEVICE***C08G 73/10* (2006.01)*C08L 79/08* (2006.01)(52) **U.S. Cl.**CPC .. *G02F 1/133753* (2013.01); *G02F 1/134309*
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(2013.01); *G02F 1/133723* (2013.01)(71) Applicant: **SHARP KABUSHIKI KAISHA,**
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(JP)(21) Appl. No.: **16/503,267**(22) Filed: **Jul. 3, 2019****Related U.S. Application Data**(60) Provisional application No. 62/702,159, filed on Jul.
23, 2018.**Publication Classification**(51) **Int. Cl.***G02F 1/1337* (2006.01)*G02F 1/1343* (2006.01)(57) **ABSTRACT**

The present invention provides a liquid crystal display device includes: a pair of substrates; a liquid crystal layer held between the substrates; an alignment film disposed on a liquid crystal layer side surface of at least one of the substrates; and a polymer layer disposed between the liquid crystal layer and the alignment film, the liquid crystal layer containing liquid crystal compounds aligned in a predetermined direction with no voltage applied, the alignment film containing a first polymer containing in its main chain at least one selected from a polyamic acid structure and a polyimide structure, the polymer layer containing a second polymer obtained by polymerizing at least one monomer containing an azobenzene group, the second polymer covalently bonded to the first polymer.



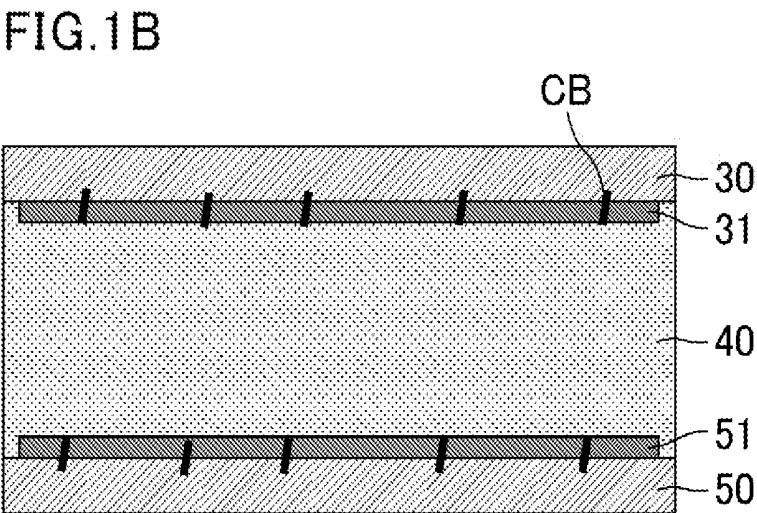
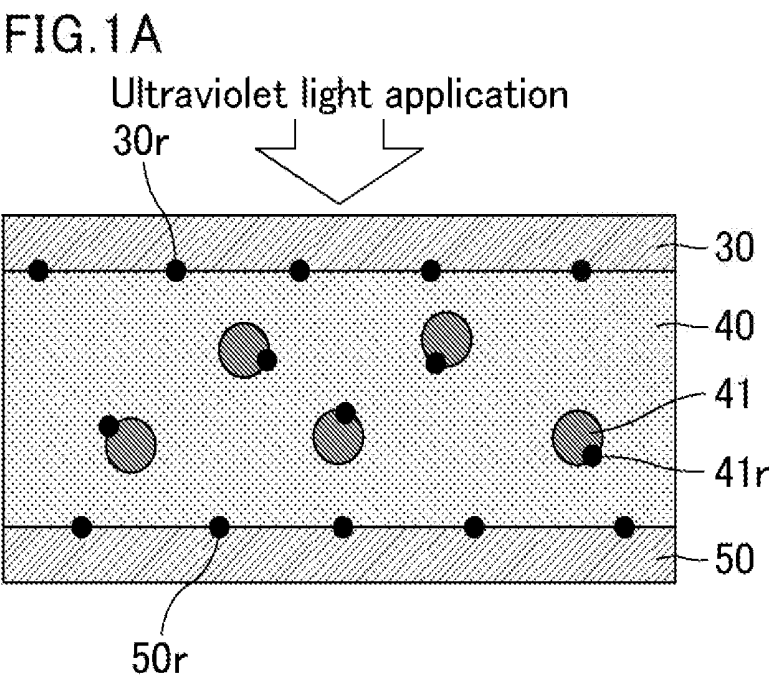


FIG. 2

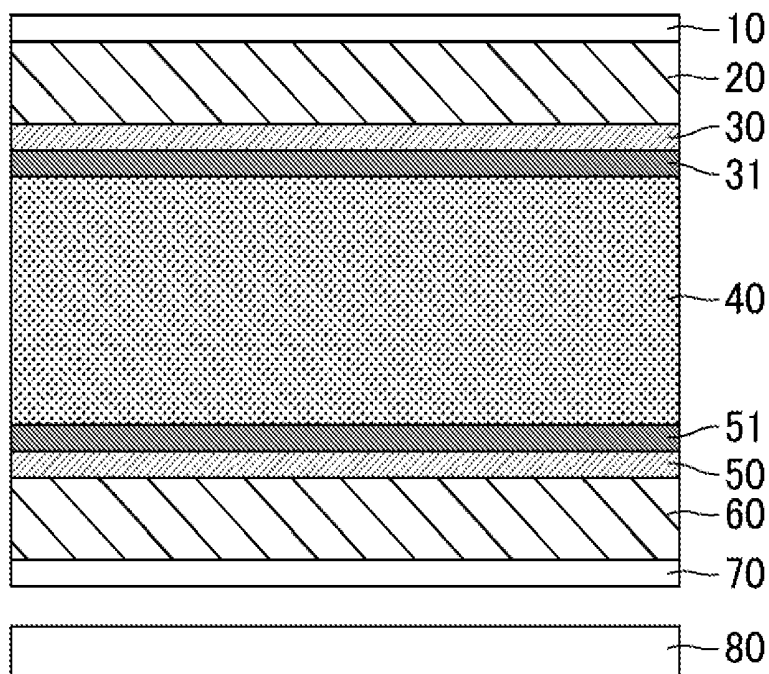
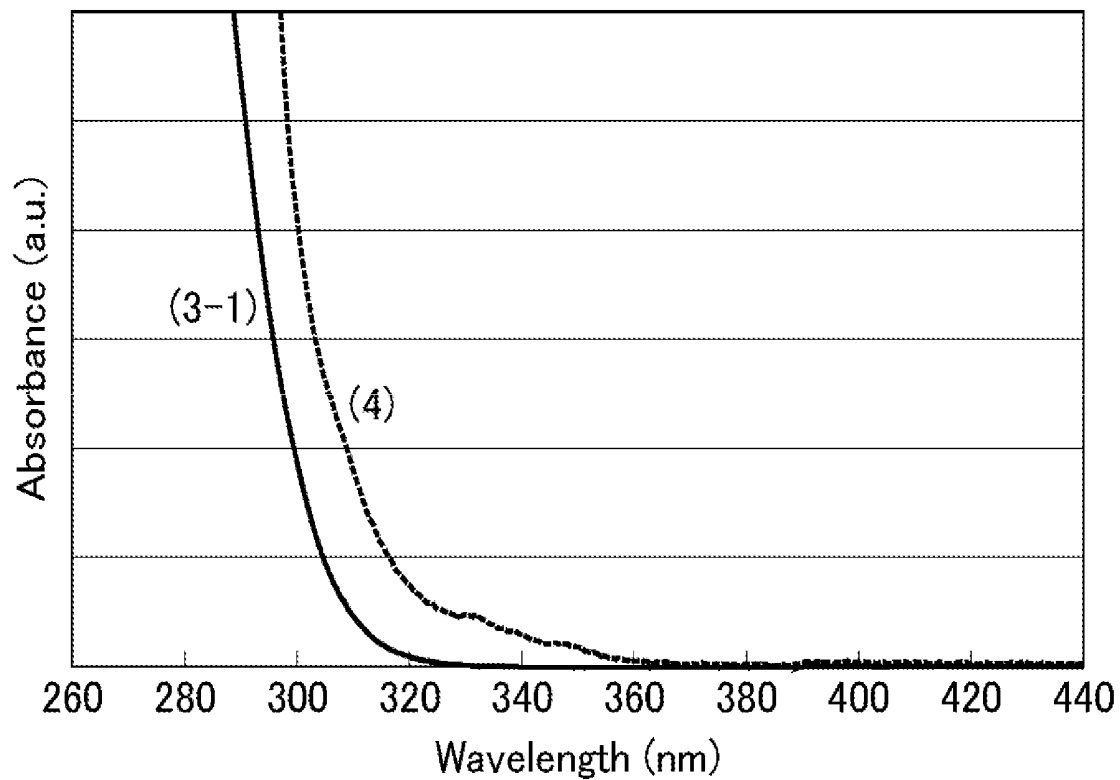


FIG. 3



LIQUID CRYSTAL DISPLAY DEVICE AND METHOD FOR MANUFACTURING LIQUID CRYSTAL DISPLAY DEVICE

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application claims priority under 35 U.S.C. § 119 to U.S. Provisional Patent Application No. 62/702,159 filed on Jul. 23, 2018, the contents of which are incorporated herein by reference in their entirety.

BACKGROUND OF THE INVENTION

Field of the Invention

[0002] The present invention relates to liquid crystal display devices and methods for manufacturing a liquid crystal display device. The present invention specifically relates to a horizontal alignment mode or vertical alignment mode liquid crystal display device including a polymer layer, and a manufacture method thereof.

Description of Related Art

[0003] Displays such as liquid crystal display devices have spread rapidly and have been used in a wide variety of applications including televisions, electronic book readers, digital photo frames, industrial appliances, personal computers (PCs), tablet PCs, smartphones, and HMDs. Liquid crystal display devices are required to have a variety of properties in these applications, and thus a variety of liquid crystal display modes have been developed.

[0004] The orientation of a liquid crystal material in a liquid crystal display device with no voltage applied is usually controlled by an alignment film on which alignment treatment has been performed. The alignment film is formed by, for example, applying an alignment film material such as a polyamic acid to a substrate and baking the material. Another technique developed to control the orientation of a liquid crystal material is a polymer sustained alignment technique (hereinafter, also referred to as the PSA technique), which polymerizes a polymerizable monomer added to a liquid crystal layer so as to form a polymer layer configured to control the orientation of a liquid crystal material on a surface of an alignment film (for example, JP 2003-177418 A, WO 2011/001579, and WO 2014/038431).

BRIEF SUMMARY OF THE INVENTION

[0005] In the case where the PSA technique is applied to a liquid crystal display device in a mode such as the in-plane switching (IPS) or fringe field switching (FFS) mode utilizing a photo-alignment technique, a vertical alignment mode, or a 4-domain reverse twisted nematic (4D-RTN) mode, a polymerization initiator cannot be used. Such a liquid crystal display device therefore causes a throughput decrease due to excessively long ultraviolet light irradiation duration and, simultaneously, the voltage holding ratio (VHR) decreases and the residual DC (rDC) voltage increases after long-term use of the liquid crystal display device due to the increased ultraviolet light irradiation dose, thereby decreasing the reliability. These undesirable changes are caused by radicals generated from polymerizable monomers remaining slightly in the liquid crystal layer.

[0006] To deal with these changes, the present inventor has studied use of a polymerization initiator with a polym-

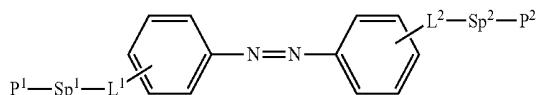
erizable group (hereinafter, also referred to as an initiator monomer). The studies, however, found that as the amount of the initiator monomer introduced into the liquid crystal material increases, the concentration of impurities from the initiator monomer increases, causing image sticking due to the VHR decrease and the residual DC voltage increase. This means that the VHR decreases and the residual DC voltage increases even when an initiator monomer or a polymerization initiator is introduced into a liquid crystal material since they remain in the liquid crystal layer, though only slightly. [0007] The same undesirable changes are presumed to occur in the liquid crystal display device disclosed in JP 2003-177418 A.

[0008] The liquid crystal display devices disclosed in WO 2011/001579 and WO 2014/038431 use a biphenyl-based monomer or a phenanthrene-based monomer, and can still be improved in terms of reduction in the VHR decrease, especially in the vertical alignment modes. These liquid crystal display devices require alignment treatment in horizontal alignment modes, and can still be improved in terms of simplification of the manufacture process.

[0009] In response to the above issues, an object of the present invention is to provide a liquid crystal display device capable of maintaining a favorable voltage holding ratio and exhibiting reduced residual DC voltage for a long period of time, and a method for manufacturing a liquid crystal display device which enables manufacture of such a liquid crystal display device.

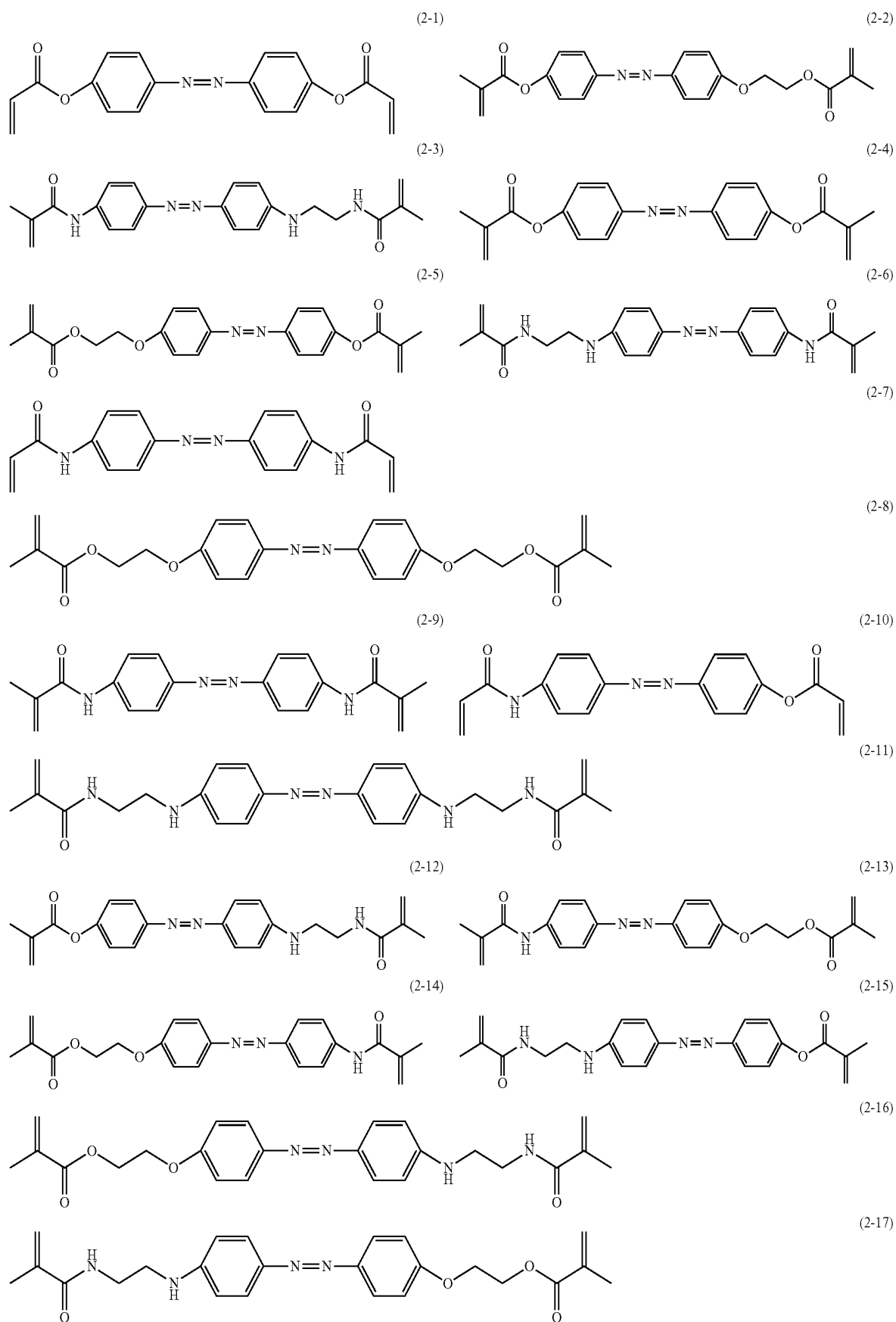
[0010] (1) One aspect of the present invention is directed to a liquid crystal display device including: a pair of substrates; a liquid crystal layer held between the substrates; an alignment film disposed on a liquid crystal layer side surface of at least one of the substrates; and a polymer layer disposed between the liquid crystal layer and the alignment film, the liquid crystal layer containing liquid crystal compounds aligned in a predetermined direction with no voltage applied, the alignment film containing a first polymer containing in its main chain at least one selected from a polyamic acid structure and a polyimide structure, the polymer layer containing a second polymer obtained by polymerizing at least one monomer including at least one monomer represented by the following formula (1), the second polymer covalently bonded to the first polymer,

(1)

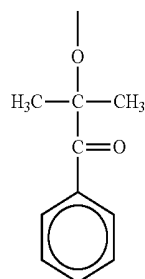


[0011] wherein P¹ and P² are the same as or different from each other, and each represent an acryloyloxy, methacryloyloxy, acryloylamino, methacryloylamino, vinyl, or vinyloxy group, Sp¹ and Sp² are the same as or different from each other, and each represent a C1-C10 linear, branched, or cyclic alkylene or alkenylene group, or a direct bond, L¹ and L² are the same as or different from each other, and each represent a —NH— group, a —N(CH₃)— group, a —O— group, a —S— group, or a direct bond, and at least one hydrogen atom in each phenylene group is optionally replaced.

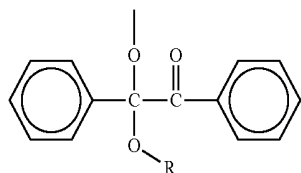
[0012] (2) In an embodiment of the present invention, the liquid crystal display device includes the structure (1), and the at least one monomer represented by the formula (1) includes at least one monomer represented by any of the following formulas (2-1) to (2-17).



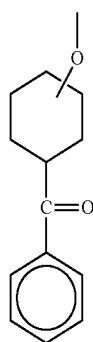
[0013] (3) In an embodiment of the present invention, the liquid crystal display device includes the structure (1) or (2), and the first polymer contains at least one functional group represented by any of the following formulas (3-1) to (3-7):



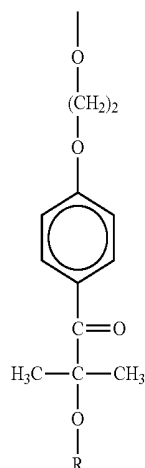
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(3-2)

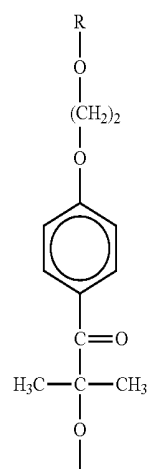


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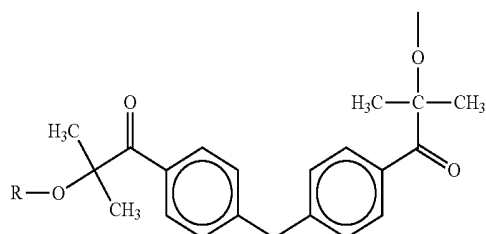


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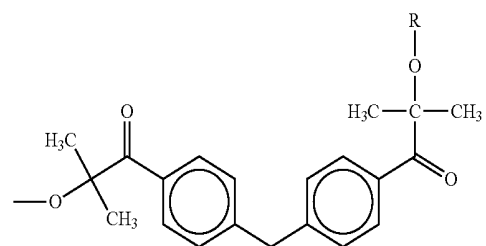
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(3-6)

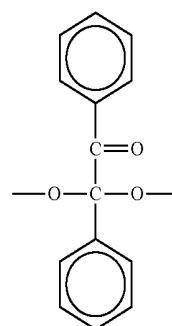


(3-7)

[0014] wherein Rs are the same as or different from each other, and each represent a C1-C40 saturated or unsaturated hydrocarbon group.

[0015] (4) In an embodiment of the present invention, the liquid crystal display device includes the structure (3), and the first polymer contains a functional group represented by the formula (3-1).

[0016] (5) In an embodiment of the present invention, the liquid crystal display device includes the structure (1), (2), (3), or (4), and the first polymer contains a functional group represented by the following formula (4).



(4)

[0017] (6) In an embodiment of the present invention, the liquid crystal display device includes the structure (1), (2), (3), (4), or (5), and the first polymer contains at least one photo-functional group.

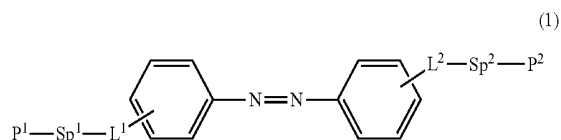
[0018] (7) In an embodiment of the present invention, the liquid crystal display device includes the structure (6), and the first polymer contains at least one photo-functional group selected from the group consisting of a cinnamate group, an azobenzene group, a chalcone group, and a coumarin group, each of which may contain a substituent.

[0019] (8) In an embodiment of the present invention, the liquid crystal display device includes the structure (1), (2), (3), (4), (5), (6), or (7), the liquid crystal display device is in a liquid crystal driving mode of a horizontal alignment mode, and the polymer layer aligns the liquid crystal compounds in a directional parallel to surfaces of the substrates.

[0020] (9) In an embodiment of the present invention, the liquid crystal display device includes the structure (1), (2), (3), (4), (5), (6), or (7), and the liquid crystal display device is in a liquid crystal driving mode of an FFS mode, an IPS mode, an MVA mode, or a 4D-RTN mode.

[0021] (10) Another aspect of the present invention is directed to a method for manufacturing a liquid crystal display device including a liquid crystal layer containing liquid crystal compounds aligned in a predetermined direction with no voltage applied, the method including: preparing a pair of substrates; forming an alignment film by applying to a surface of at least one of the substrates an alignment agent that contains a first polymer containing at least one selected from a polyamic acid structure and a polyimide structure in its main chain; forming a liquid crystal layer by sealing, between the substrates on at least one of which the alignment film is formed, a liquid crystal composition containing a liquid crystal material and at least

one monomer including at least one monomer represented by the following formula (1); and forming a polymer layer from a second polymer between the alignment film and the liquid crystal layer by irradiating the liquid crystal layer with ultraviolet light, the second polymer being obtained by polymerizing the at least one monomer including at least one monomer represented by the following formula (1), the forming a polymer layer including covalently bonding the second polymer to the first polymer,



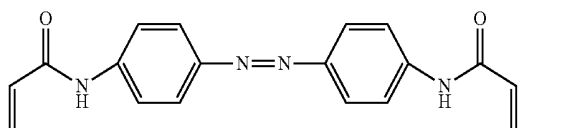
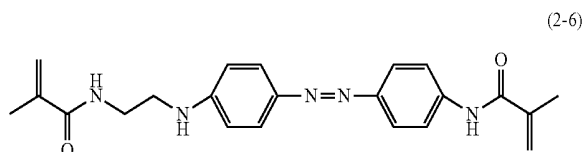
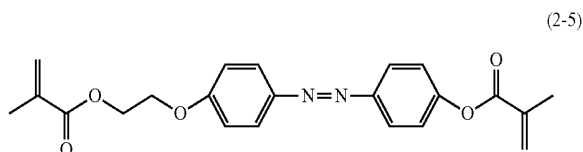
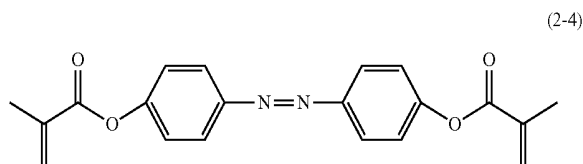
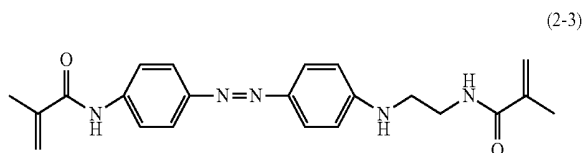
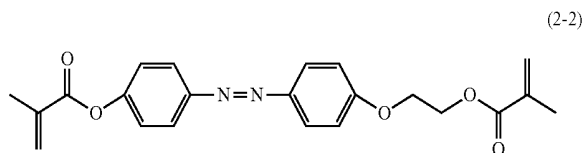
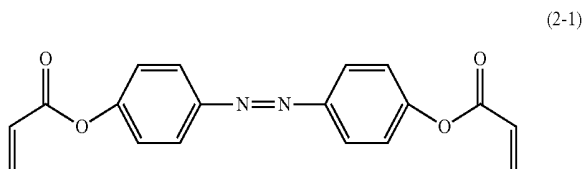
[0022] wherein P^1 and P^2 are the same as or different from each other, and each represent an acryloyloxy, methacryloyloxy, acryloylamino, methacryloylamino, vinyl, or vinyloxy group,

[0023] Sp^1 and Sp^2 are the same as or different from each other, and each represent a C1-C10 linear, branched, or cyclic alkylene or alkenylene group, or a direct bond,

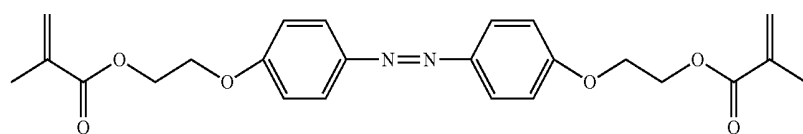
[0024] L^1 and L^2 are the same as or different from each other, and each represent a $-\text{NH}-$ group, a $-\text{N}(\text{CH}_3)-$ group, a $-\text{O}-$ group, a $-\text{S}-$ group, or a direct bond, and

[0025] at least one hydrogen atom in each phenylene group is optionally replaced.

[0026] (11) In an embodiment of the present invention, the method includes the process (10), and the at least one monomer represented by the formula (1) includes at least one monomer represented by any of the following formulas (2-1) to (2-17).



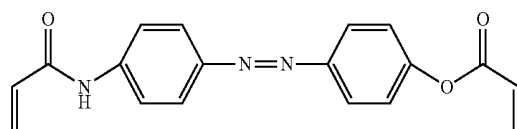
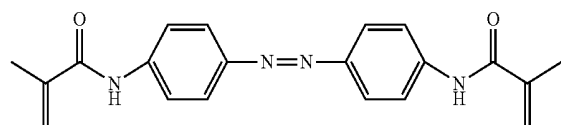
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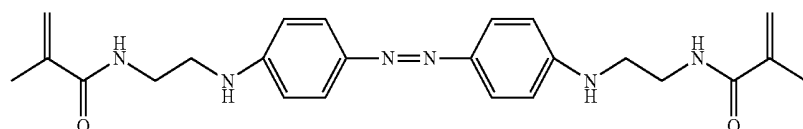
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(2-10)

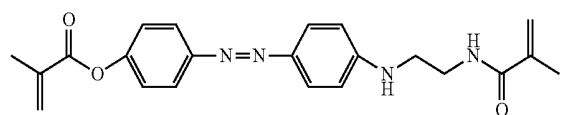


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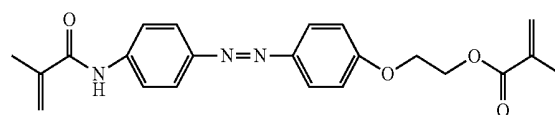


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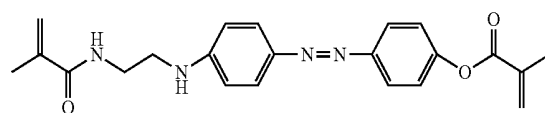
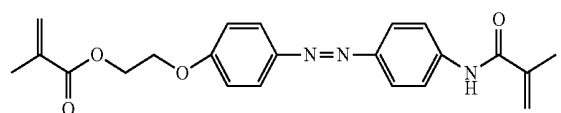
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(2-14)



(2-15)

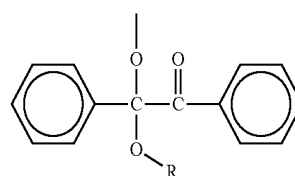


(2-16)

(2-17)

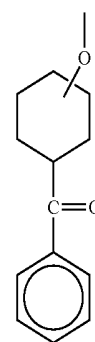
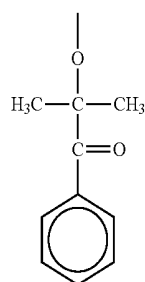
[0027] (12) In an embodiment of the present invention, the method includes the process (10) or (11), and the first polymer contains at least one functional group represented by any of the following formulas (3-1) to (3-7):

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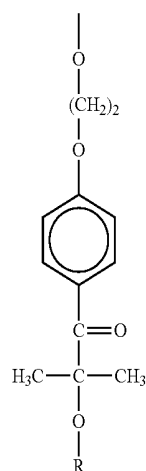
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(3-1)



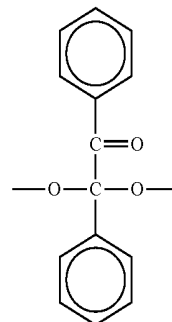
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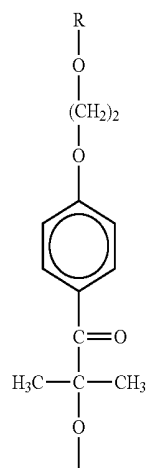


(3-4)

[0030] (14) In an embodiment of the present invention, the method includes the process (10), (11), (12), or (13), and the first polymer contains a functional group represented by the following formula (4).



(4)



(3-5)

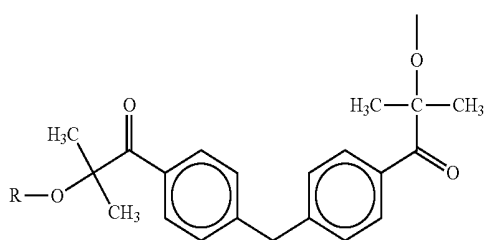
[0031] (15) In an embodiment of the present invention, the method includes the process (10), (11), (12), (13), or (14), and the first polymer contains at least one photo-functional group.

[0032] (16) In an embodiment of the present invention, the method includes the process (10), (11), (12), (13), (14), or (15), and the forming a polymer layer includes irradiating the liquid crystal layer with polarized ultraviolet light so as to polymerize the at least one monomer including at least one monomer represented by the formula (1).

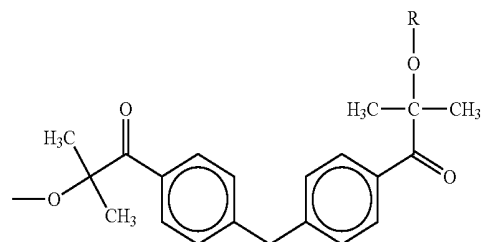
[0033] (17) In an embodiment of the present invention, the method includes the process (10), (11), (12), (13), (14), or (15), and the forming a polymer layer includes irradiating the liquid crystal layer with unpolarized ultraviolet light so as to polymerize the at least one monomer including at least one monomer represented by the formula (1).

[0034] JP 2003-177418 A mentions a polymerization initiator, but discloses only a method including adding a polymerization initiator to a liquid crystal layer. JP 2003-177418 A fails to examine covalent bonding of the polymer of the polymer layer and the polymer of the alignment film. WO 2011/001579 and WO 2014/038431 fail to examine the monomer represented by the formula (1).

[0035] The present invention achieves a liquid crystal display device capable of maintaining a favorable voltage holding ratio and reduced residual DC voltage for a long period of time.



(3-6)



(3-7)

[0028] wherein Rs are the same as or different from each other, and each represent a C1-C40 saturated or unsaturated hydrocarbon group.

[0029] (13) In an embodiment of the present invention, the method includes the process (12), and the first polymer contains a functional group represented by the formula (3-1).

BRIEF DESCRIPTION OF THE DRAWINGS

[0036] FIG. 1A is a schematic cross-sectional view of a liquid crystal cell before polymerization of monomers, illustrating polymer layer formation processes in a method for manufacturing a liquid crystal display device of Embodiment 1.

[0037] FIG. 1B is a schematic cross-sectional view of the liquid crystal cell after the polymerization of monomers, illustrating the polymer layer formation processes in the method for manufacturing the liquid crystal display device of Embodiment 1.

[0038] FIG. 2 is a schematic cross-sectional view of the liquid crystal display device of Embodiment 1.

[0039] FIG. 3 is a graph of absorption spectra of a functional group represented by the below formula (3-1) and a moiety excluding a hydrocarbon group of a functional group

represented by the below formula (3-2) (functional group represented by the below formula (4)).

DETAILED DESCRIPTION OF THE INVENTION

[0040] The present invention is described in more detail below based on the following embodiment with reference to the drawing. The embodiment, however, is not intended to limit the scope of the present invention. The configurations of the embodiment may appropriately be combined or modified within the spirit of the present invention.

[0041] The “viewing surface side” as used herein means the side closer to the screen (display surface) of the display device. The “back surface side” means the side farther from the screen (display surface) of the display device. The “room temperature” is 15° C. or higher and 40° C. or lower, unless otherwise specified.

[0042] The “photo-functional group” as used herein means a functional group that can undergo a photoreaction. The photo-functional group preferably can undergo a structural change such as dimerization (formation of dimers), isomerization, photo-Fries rearrangement, and decomposition (cleavage) when irradiated with light (electromagnetic waves) such as ultraviolet rays or visible light (preferably polarized light, more preferably polarized ultraviolet rays, particularly preferably linearly polarized ultraviolet rays), and thereby exhibit an ability of controlling the alignment of liquid crystal compounds. Specific examples of the photo-functional group include an azobenzene group, a chalcone group, a cinnamate group, a coumarin group, a tolane group, a stilbene group, and a cyclobutane ring.

[0043] The nematic-isotropic phase transition temperature (T_{ni}) as used herein is determined by visually observing the liquid crystal state or the isotropic state while varying the temperature using a device such as one from Mettler. The temperature can also be determined by a technique of determining the temperature at which phase transition occurs using a differential scanning calorimeter (DSC).

[0044] The mode herein in which liquid crystal compounds (molecules) are aligned in a direction substantially parallel to a main surface of each of the pair of substrates with no voltage applied is also referred to as a horizontal alignment mode. The expression “substantially parallel” means that, for example, the pre-tilt angle of liquid crystal compounds is 0° or greater and 5° or smaller from the main surface of each substrate. The mode in which liquid crystal compounds are aligned in a direction substantially perpendicular to the main surface of each of the pair of substrates with no voltage applied is also referred to as a vertical alignment mode. The expression “substantially perpendicular” means that, for example, the pre-tilt angle of liquid crystal compounds is 85° or greater and 90° or smaller from the main surface of the substrate. The pre-tilt angle is an angle of the major axis of a liquid crystal material (liquid crystal compound) from a surface of a substrate when the voltage applied to the liquid crystal layer is lower than the threshold voltage (including the case of no voltage application), with the substrate surface taken as 0° and the line normal to the substrate as 90°. The present invention is applicable to both horizontal alignment mode liquid crystal display devices and vertical alignment mode liquid crystal display devices.

Embodiment 1

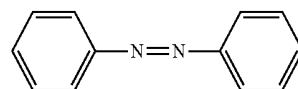
[0045] The present embodiment is summarized first. The present embodiment takes the following measures (1) and (2) to overcome the above issues.

[0046] (1) A polymer constituting an alignment film is chemically modified with (covalently bonded to) a functional group capable of initiating radical polymerization under light (hereinafter, also referred to as an initiator functional group).

[0047] In this measure, a polymerization initiator or an initiator monomer is not introduced into the liquid crystal layer, and the moiety functioning as a polymerization initiator can be immobilized on an alignment film surface. This prevents the polymerization initiator components from remaining in the liquid crystal layer, reducing generation of image sticking due to polymerization initiator components. Also, just adding a low molecular weight polymerization initiator into an alignment film may not be enough to prevent the polymerization initiator, which has a low molecular weight, from dissolving in the liquid crystal layer. The point is to introduce the initiator functional group into the polymer (preferably a side chain thereof) constituting the alignment film and covalently bond the polymer to the polymer layer.

[0048] (2) A bifunctional group (monomer having two polymerizable groups) containing an azobenzene group (functional group derived from azobenzene) is introduced into the liquid crystal layer.

[0049] Since azobenzene groups become alignable when irradiated with polarized ultraviolet light unlike conventional monomers for PSA, a polymer layer (PSA layer) capable of horizontally aligning liquid crystal compounds when irradiated with polarized ultraviolet light can be formed (this process requires heating at the T_{ni} of the liquid crystal material or higher) without alignment control provided by the alignment film. This can overcome the issue of misalignment in bonding of a pair of substrates. Azobenzene is a chemical structure represented by the following formula (A). A monomer containing an azobenzene group contains an azo group ($—N=N—$) and causes interaction between an unpaired electron of the azobenzene group and an unpaired electron of a polyimide (acid anhydride). A polymer of the monomer containing an azobenzene group therefore easily separates from the liquid crystal layer to form a polymer layer.



(A)

[0050] FIG. 1A is a schematic cross-sectional view of a liquid crystal cell before polymerization of monomers, illustrating polymer layer formation processes in a method for manufacturing a liquid crystal display device of Embodiment 1. FIG. 1B is a schematic cross-sectional view of the liquid crystal cell after the polymerization of monomers, illustrating the polymer layer formation processes in the method for manufacturing the liquid crystal display device of Embodiment 1. As shown in FIG. 1A, application of ultraviolet light not only generates radicals **41r** in monomers **41** but also generates radicals **30r** and **50r** in alignment films **30** and **50** (preferably, their polymer side chains), respec-

tively. As shown in FIG. 13, the monomers **41** in the liquid crystal layer **40** are covalently bonded to the alignment films **30** and **50** (preferably, their polymer side chains). This enables the design of a PSA liquid crystal display device in which the alignment films **30** and **50** are bonded to the polymer layers **31** and **51** by covalent bonds CB, respectively. The covalent bonds between the alignment film **30** and the corresponding polymer layer **31** and between the alignment film **50** and the corresponding polymer layer **51** increase the polymerization rate of the monomers **41** to increase the formation rate of the polymer layer, leading to a favorable VHR and reduced rDC voltage in long-term use. With the monomers **41** containing an azobenzene group in the liquid crystal layer **40**, the polymer layers **31** and **51** capable of achieving the horizontal alignment can be formed by applying polarized ultraviolet light.

[0051] JP 2003-177418 A fails to disclose or suggest such features of the present embodiment. The liquid crystal display devices disclosed in WO 2011/001579 and WO 2014/038431 use a biphenyl-based monomer or a phenanthrene-based monomer as described above. In a horizontal alignment mode, biphenyl-based monomers and phenanthrene-based monomers are insufficient in enhancing the stability of the alignment azimuth of liquid crystal monomers. A monomer containing an azobenzene group, which highly absorbs polarized light, is suitable. Use of a monomer containing an azobenzene group improves the horizontal alignment state of liquid crystal molecules, achieves a high contrast ratio, and enhances the alignment stability of liquid crystal molecules, thereby reducing the chances of a contrast ratio decrease in long-term use of the liquid crystal display device.

[0052] FIG. 2 is a schematic cross-sectional view of a liquid crystal display device of Embodiment 1. As shown in FIG. 2, the liquid crystal display device of Embodiment 1 is a horizontal alignment mode or vertical alignment mode liquid crystal display device including, in the following order from the viewing surface side to the back surface side, a first linear polarizer **10**, a counter substrate **20**, an alignment film **30**, a polymer layer **31**, a liquid crystal layer **40**, a polymer layer **51**, an alignment film **50**, a thin-film transistor (TFT) substrate **60**, a second linear polarizer **70**, and a backlight **80**. The polymer layers **31** and **51** may be disposed on the entire or part (in either case, within the region surrounded by the sealant in a plan view) of the liquid crystal layer **40** side surfaces of the alignment films **30** and **50**, respectively.

[0053] The first linear polarizer **10** can be, for example, a polarizer (absorptive polarizer) obtained by dyeing a polyvinyl alcohol (PVA) film with an anisotropic material such as an iodine complex (or dye) to adsorb the anisotropic material on the polyvinyl alcohol film, and stretching the film for alignment. Typically, each surface of a PVA film is laminated with a protective film such as a triacetyl cellulose (TAC) film in practical use for sufficient mechanical strength and sufficient moisture and heat resistance.

[0054] The counter substrate **20** is a color filter (CF) substrate that includes, in the following order from the viewing surface side to the back surface side, a transparent substrate (not illustrated), color filters/black matrix (not illustrated), and a flattening film as necessary.

[0055] The transparent substrate may be, for example, a glass substrate or a plastic substrate.

[0056] The color filters/black matrix have a structure in which red color filters, green color filters, and blue color filters are arranged in a plane and partitioned by a black matrix. The red color filters, the green color filters, the blue color filters, and the black matrix are each made of, for example, a transparent resin containing a pigment. Typically, a combination of a red color filter, a green color filter, and a blue color filter is arranged in each pixel, and the desired color is achieved in each pixel by mixing colors of the red color filter, the green color filter, and the blue color filter while controlling the amount of light passing through the filters.

[0057] The alignment films **30** and **50** may be horizontal alignment films configured to align liquid crystal compounds in a direction substantially parallel to their surfaces or may be vertical alignment films configured to align liquid crystal compounds in a direction substantially perpendicular to their surfaces. The alignment films **30** and **50** may be photo-alignment films containing a photo-functional group and having been subjected to photo-alignment as the alignment treatment, rubbed alignment films having been subjected to rubbing as the alignment treatment, or alignment films not having been subjected to any alignment treatment.

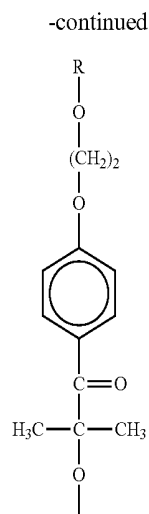
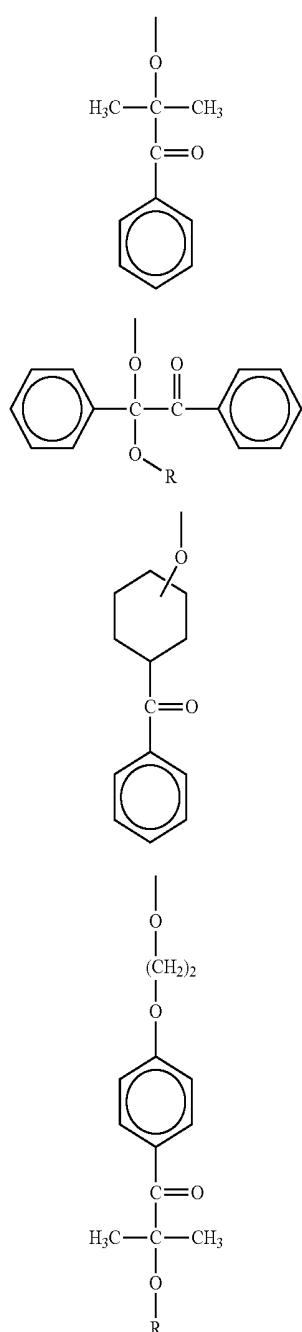
[0058] Each of the alignment films **30** and **50** contains a first polymer containing in its main chain at least one selected from a polyamic acid structure and a polyimide structure (hereinafter, also referred to as a polyimide-based first polymer).

[0059] The polyimide-based first polymer has a diamine-derived structure and a tetracarboxylic dianhydride-derived structure as repeating structures, and is obtained by polymerizing at least one diamine and at least one tetracarboxylic dianhydride.

[0060] The polyimide-based first polymer is covalently bonded to the later-described second polymer contained in each of the polymer layers **31** and **51**. This can increase the formation rates of the polymer layers **31** and **51** (polymerization rate of the monomers), reducing the ultraviolet light irradiation dose for the liquid crystal layer in the polymer layer formation. These can prevent a VHR decrease and a residual DC voltage increase even after long-term use of the liquid crystal display device of the present embodiment.

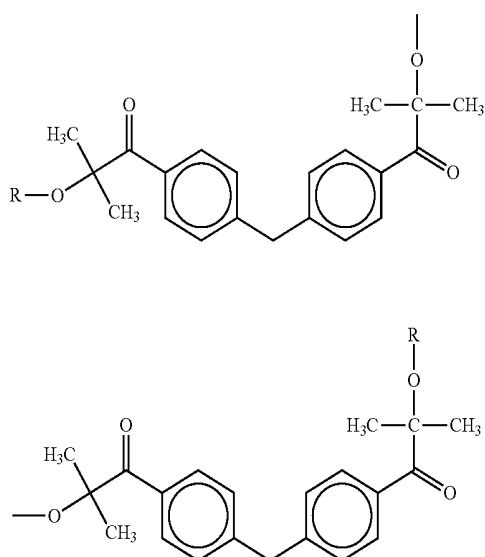
[0061] The polyimide-based first polymer preferably has at least one functional group selected from those represented by any of the following formulas (3-1) to (3-7). These functional groups are initiator functional groups and thus can generate radicals when irradiated with ultraviolet light, thereby being functionable as a radical polymerization initiator. Meanwhile, these functional groups, constituting the polyimide-based first polymer, hardly dissolve in the liquid crystal layer **40**, unlike common polymerization initiators and initiator monomers to be added to the liquid crystal layer in the PSA technique. Also, radicals generated from these functional groups react with monomers added to the liquid crystal layer **40**, and the alignment films **30** and **50** form

covalent bonds with polymers (the later-described second polymer) constituting the polymer layers **31** and **51**, respectively. This, as described above, can increase the formation rates of the polymer layers **31** and **51** (polymerization rate of the monomers), reducing the ultraviolet light irradiation dose for the liquid crystal layer **40** in the polymer layer formation. These can more effectively prevent a VHR decrease and a residual DC voltage increase even after long-term use of the liquid crystal display device of the present embodiment. The initiator functional groups are preferably introduced into a side chain of the polyimide-based first polymer.



-continued

(3-5)



(3-6)

(3-7)

(3-3)

(3-4)

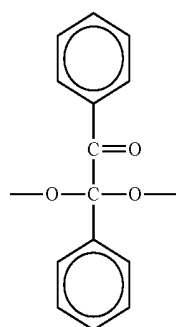
[0062] In the formulas, Rs are the same as or different from each other, and each represent a C1-C40 saturated or unsaturated hydrocarbon group.

[0063] FIG. 3 is a graph of absorption spectra of a functional group represented by the below formula (3-1) and a moiety excluding a hydrocarbon group of a functional group represented by the below formula (3-2) (functional group represented by the below formula (4)). As shown in FIG. 3, the functional group represented by the formula (3-1) has a shorter absorption wavelength than the functional group represented by the formula (3-2). For this reason, when an unreacted initiator functional group remains and the liquid crystal display device of the present embodiment is used for a long period of time, i.e., several years, the functional group represented by the formula (3-2) is likely to cleave under the backlight illumination or external light, generating radicals. In the case of a photocleavage under the backlight illumination, radical generation is likely to start from the alignment film **50**, which is closer to the backlight **80**. In the case of a photocleavage under external light, radical generation is likely to start from the alignment film **30**, which is closer to

the external light (i.e., the side remote from the backlight **80**). Such photocleavages are presumed to cause a gradual increase in residual DC voltage. Meanwhile, the functional group represented by the formula (3-1) is not likely to cause asymmetrization due to the above radical generation even when the liquid crystal display device of the present embodiment is used for a very long period of time. The point of the present embodiment is as follows. Even when unreacted initiator functional groups are present, they are not in the liquid crystal layer **40** (since they are chemically bonded to the alignment films **30** and **50**), and thus the reliability is not decreased in normal practical use. Yet, in consideration of cases where the display device is used for a very long period of time and the backlight **80** used is a high-luminance backlight, the functional group represented by the formula (3-1) is preferred to the functional group represented by the formula (3-2). A functional group represented by any of the formulas (3-3) to (3-7) has a short absorption wavelength as with the functional group represented by the formula (3-1), and is therefore preferred to the functional group represented by the formula (3-2) in terms of the very long-term reliability.

[0064] When the functional group represented by any of the formulas (3-1) and (3-3) to (3-7) is introduced into a side chain of the polyimide-based first polymer, the binding site (—O— group) with the side chain or main chain and the phenyl group can be positioned linearly, so that the side chain containing any of these functional groups can be prevented from becoming bulky. This technique therefore eliminates the need to consider the influence on introduction and the introduction ratio of another side chain (e.g., vertical alignment functional group, photo-functional group). In contrast, when the functional group represented by the formula (3-2) is introduced into a side chain of the polyimide-based first polymer, the binding site (—O— group) with the side chain or main chain and the two phenyl groups form a bent shape (become perpendicular to each other), so that the side chain containing the functional group represented by the formula (3-2) becomes bulky. This may inhibit introduction of another side chain (e.g., vertical alignment functional group, photo-functional group). In the case where another side chain is difficult to introduce due to its chemical structure, the molecular structure design of the alignment films **30** and **50** may be limited.

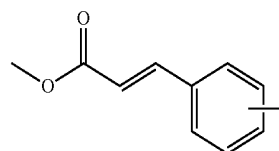
[0065] In particular, the polyimide-based first polymer preferably has a functional group represented by the formula (3-1) and preferably has a functional group represented by the following formula (4).



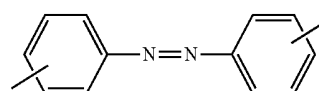
(4)

[0066] The polyimide-based first polymer preferably has at least one photo-functional group, more preferably at least one photo-functional group selected from the group consisting of a cinnamate group, an azobenzene group, a chalcone group, and a coumarin group, each of which may contain a substituent.

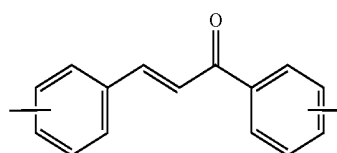
[0067] Preferred examples of the substituent include, but are not limited to, a halogen group, a methyl group, a methoxy group, an ethyl group, and an ethoxy group. These may be used alone or in combination with each other. In other words, the substituent preferably includes at least one substituent selected from the group consisting of a halogen group, a methyl group, a methoxy group, an ethyl group, and an ethoxy group. The halogen group is preferably a fluoro group or a chloro group. In the case where the photo-functional group contains a substituent, the substituent usually replaces at least one hydrogen atom in a ring structure, such as a phenylene group, of the photo-functional group. The photo-functional group may be a monovalent functional group, but is preferably a divalent cinnamate group represented by the following formula (B-1), a divalent azobenzene group represented by the following formula (B-2), a divalent chalcone group represented by the following formula (B-3), or a divalent coumarin group represented by the following formula (B-4).



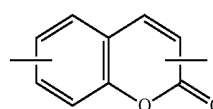
(B-1)



(B-2)

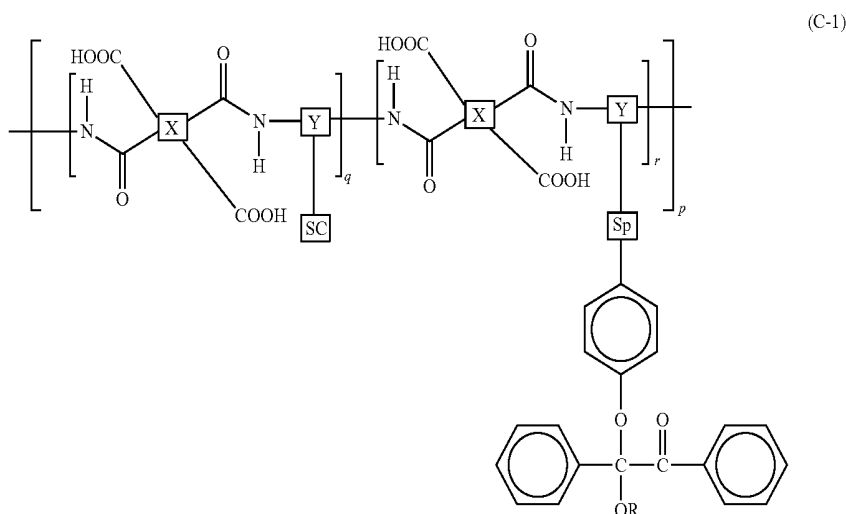


(B-3)



(B-4)

[0068] Preferred examples of the polyimide-based first polymer include polyamic acid structures represented by the following formula (C-1) or (C-2) (the polyamic acid structures may be at least partially imidized into polyimide structures). These may be used alone or in combination with each other.

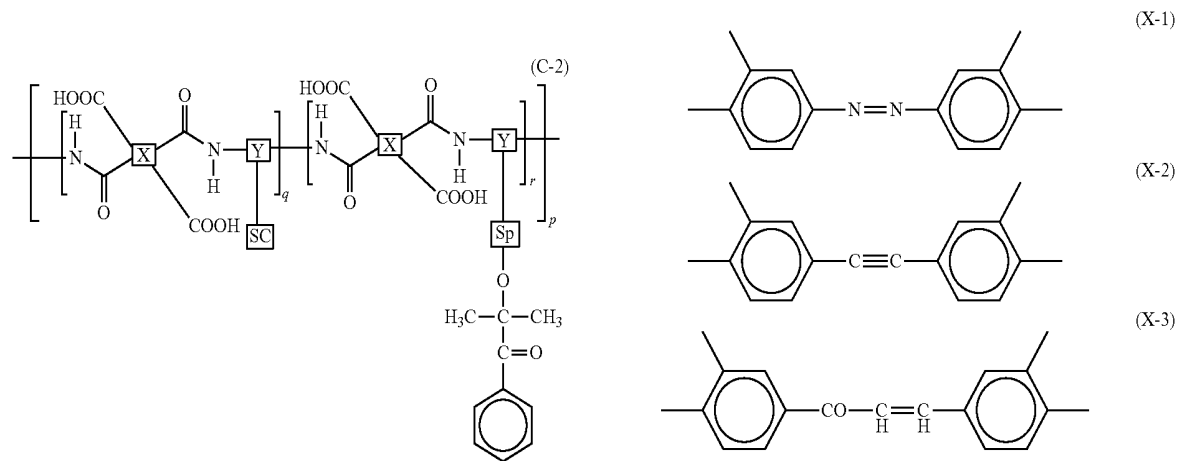


[0069] In the formula, X represents a tetravalent organic group, Y represents a trivalent organic group, SC represents a side chain, Sp represents a spacer or a direct bond, R represents a monovalent organic group, p represents a degree of polymerization, p, q, and r are each independently an integer of 1 or greater, p is preferably 10 or greater, and q and r satisfy the relation $0 < r/(q+r) \leq 1$.

[0070] Specific examples of R in the formula (C-1) include methyl, ethyl, propyl, isopropyl, butyl, and isobutyl groups. These may be used alone or in combination with each other.

mode), q and r in the formulas (C-1) and (C-2) are not limited as long as they satisfy the relation $0 < r/(q+r) \leq 1$.

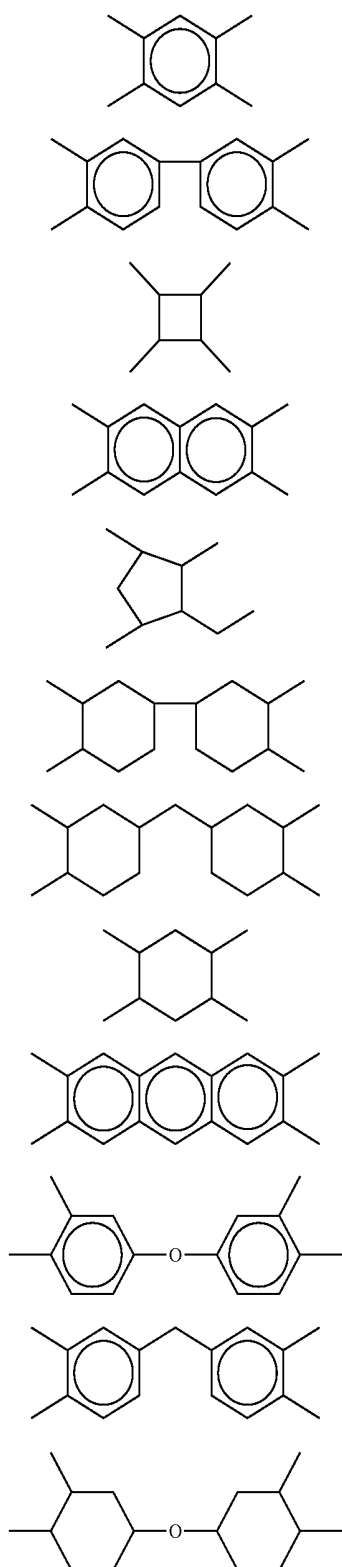
[0073] In the formulas (C-1) and (C-2), in the case where X contains a photo-functional group, X may be, for example, a group represented by any of the following formulas (X-1) to (X-4). These groups can be used both in the case where the alignment films **30** and **50** are horizontal alignment films and in the case where the alignment films **30** and **50** are vertical alignment films. These groups may be used alone or in combination with each other,



[0071] In the formula, X represents a tetravalent organic group, Y represents a trivalent organic group, SC represents a side chain, Sp represents a spacer or a direct bond, p represents a degree of polymerization, p, q, and r are each independently an integer of 1 or greater, p is preferably 10 or greater, and q and r satisfy the relation $0 < r/(q+r) \leq 1$.

[0072] In the case where the alignment films **30** and **50** are vertical alignment films (in the vertical alignment mode), q and r in the formulas (C-1) and (C-2) preferably satisfy the relation $0 < r/(q+r) \leq 0.5$, more preferably the relation $0 < r/(q+r) \leq 0.4$. In the case where the alignment films **30** and **50** are horizontal alignment films (in the horizontal alignment

[0074] In the formulas (C-1) and (C-2), in the case where X contains no photo-functional group, X may be, for example, a group represented by any of the following formulas (X-5) to (X-16). These groups can be used both in the case where the alignment films **30** and **50** are horizontal alignment films and in the case where the alignment films **30** and **50** are vertical alignment films. These groups may be used alone or in combination with each other.



[0075] In the formulas (C-1) and (C-2), in the case where Y contains a photo-functional group, Y may be, for example, a group represented by any of the following formulas (Y-1) to (Y-8). These groups can be used both in the case where the alignment films **30** and **50** are horizontal alignment films and

in the case where the alignment films **30** and **50** are vertical alignment films. These groups may be used alone or in combination with each other.

(X-5)

(X-6)

(X-7)

(X-8)

(X-9)

(X-10)

(X-11)

(X-12)

(X-13)

(X-14)

(X-15)

(X-16)

(Y-1)

(Y-2)

(Y-3)

(Y-4)

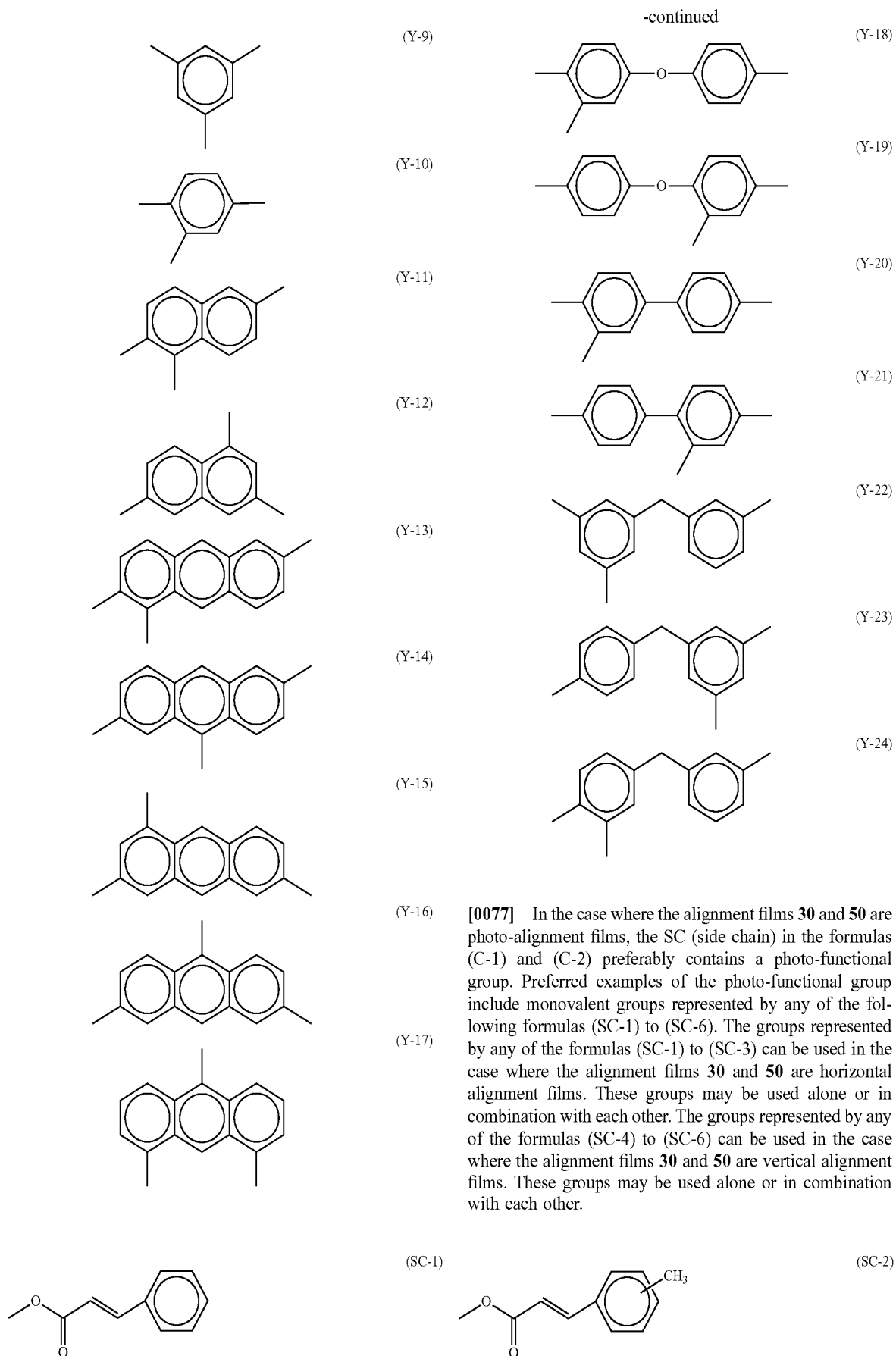
(Y-5)

(Y-6)

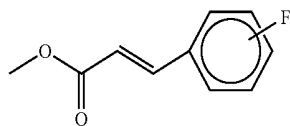
(Y-7)

(Y-8)

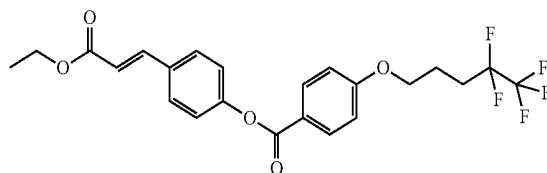
[0076] In the formulas (C-1) and (C-2), in the case where Y contains no photo-functional group, Y may be, for example, a group represented by any of the following formulas (Y-9) to (Y-24). These groups can be used both in the case where the alignment films **30** and **50** are horizontal alignment films and in the case where the alignment films **30** and **50** are vertical alignment films. These groups may be used alone or in combination with each other.



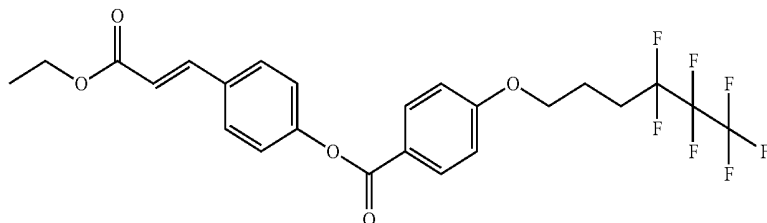
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(SC-3)



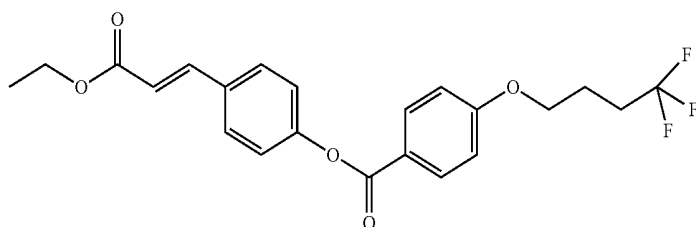
(SC-4)



(SC-5)



(SC-6)



[0078] In the case where the alignment films **30** and **50** are not photo-alignment films, the SC (side chain) in the formulas (C-1) and (C-2) may contain a horizontal alignment functional group other than a photo-functional group, such as a monovalent group represented by any of the following formulas (SC-7) to (SC-13). Also, the SC (side chain) may be excluded and a hydrogen atom (hydrogen group) may be bonded to Y. These groups can be used in the case where the alignment films **30** and **50** are horizontal alignment films. These groups may be used alone or in combination with each other.

[0079] In the case where the alignment films **30** and **50** are not photo-alignment films, the SC (side chain) in the formulas (C-1) and (C-2) may contain a vertical alignment functional group other than a photo-functional group, such as a monovalent group represented by any of the following formulas (SC-14) to (SC-20). These groups can be used in the case where the alignment films **30** and **50** are vertical alignment films. These groups may be used alone or in combination with each other.



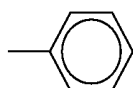
(SC-7)



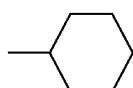
(SC-8)



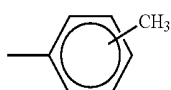
(SC-9)



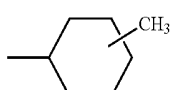
(SC-10)



(SC-11)

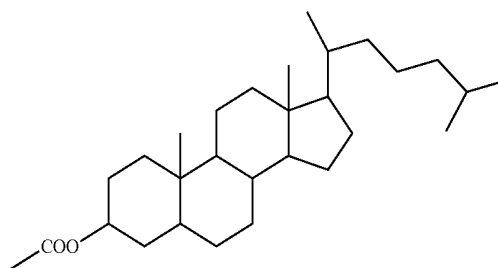


(SC-12)

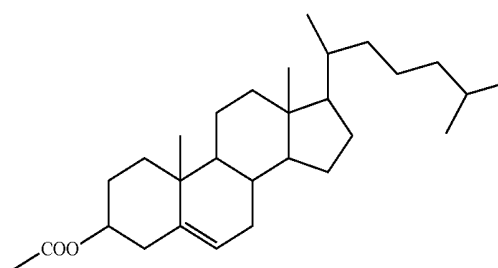


(SC-13)

(SC-14)

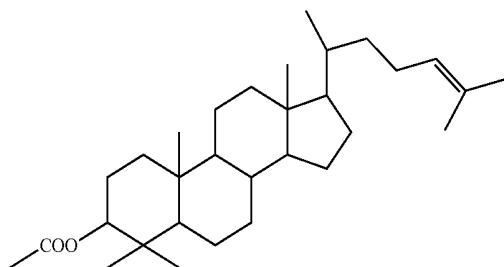


(SC-15)

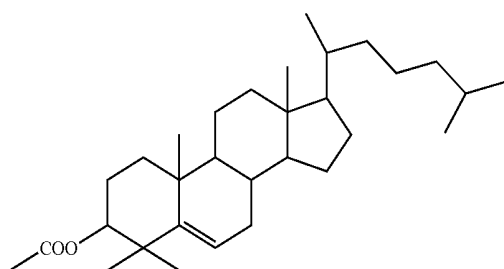


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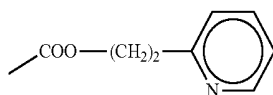
(SC-16)



(SC-17)

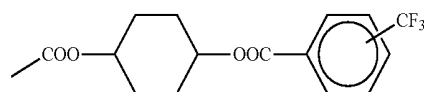


(SC-18)

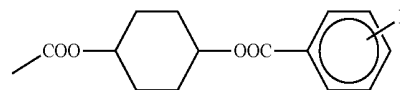


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(SC-19)



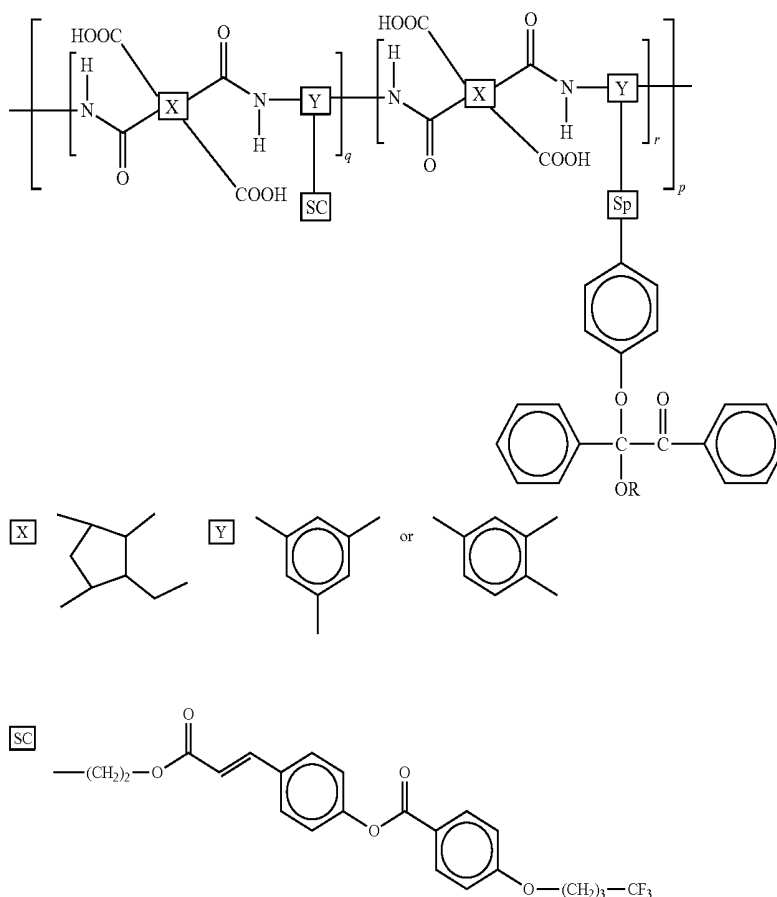
(SC-20)



[0080] The polyamic acid structure represented by the formula (C-2) (the polyamic acid structure may be at least partially imidized into a polyimide structure) contains an initiator functional group represented by the formula (3-1), but may contain at least one initiator functional group represented by any of the formulas (3-2) to (3-7) instead of the at least one initiator functional group represented by the formula (3-1).

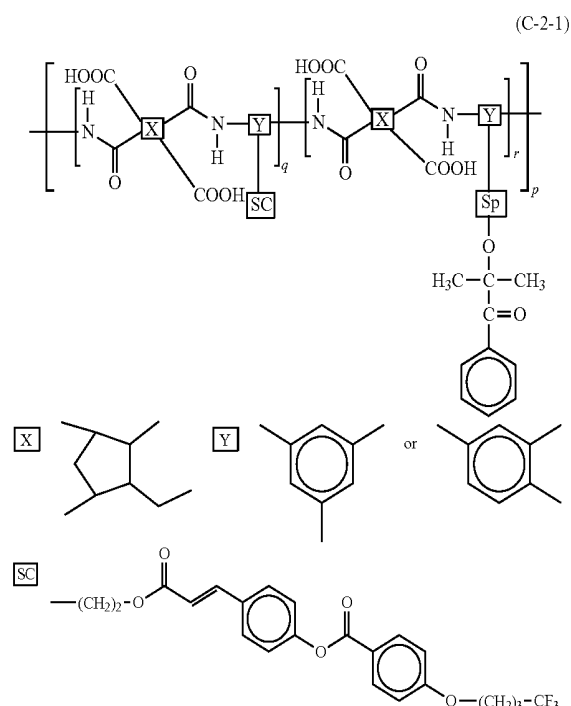
[0081] Particularly preferred examples of the polyimide-based first polymer include polyamic acid structures represented by any of the following formulas (C-1-1) and (C-2-1) (the polyamic acid structures may be at least partially imidized into polyimide structures). These may be used alone or in combination with each other.

(C-1-1)



[0082] In the formula, Sp represents a spacer or a direct bond, p represents a degree of polymerization, p, q, and r are each independently an integer of 1 or greater, p is preferably 10 or greater, and q and r satisfy the relation $0 < r/(q+r) \leq 1$ (in the case of the vertical alignment mode, preferably the relation $0 < r/(q+r) \leq 0.5$, more preferably the relation $0 < r/(q+r) \leq 0.4$).

[0083] Specific examples of R in the formula (C-1-1) include methyl, ethyl, propyl, isopropyl, butyl, and isobutyl groups. These may be used alone or in combination with each other.



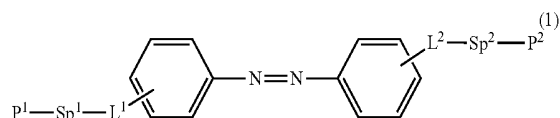
[0084] In the formula, Sp represents a spacer or a direct bond, p represents a degree of polymerization, p, q, and r are each independently an integer of 1 or greater, p is preferably 10 or greater, and q and r satisfy the relation $0 < r/(q+r) \leq 1$ (in the case of the vertical alignment mode, preferably the relation $0 < r/(q+r) \leq 0.5$, more preferably the relation $0 < r/(q+r) \leq 0.4$).

[0085] The polyamic acid structure represented by the 1.0 formula (C-2-1) (the polyamic acid structure may be at least partially imidized into a polyimide structure) contains an initiator functional group represented by the formula (3-1), but may contain at least one initiator functional group represented by any of the formulas (3-2) to (3-7) instead of the at least one initiator functional group represented by the formula (3-1).

[0086] For a high contrast ratio, the alignment films 30 and 50 are preferably photo-alignment films. In the case where the alignment films 30 and 50 each contain a structure represented by the formula (C-1) and/or the formula (C-2), the alignment films 30 and 50 are made into photo-alignment films by introducing a photo-functional group into at least one selected from X, Y, and SC (side chain).

[0087] Each of the polymer layers 31 and 51 contains a second polymer obtained by polymerizing at least one monomer including at least one monomer represented by the

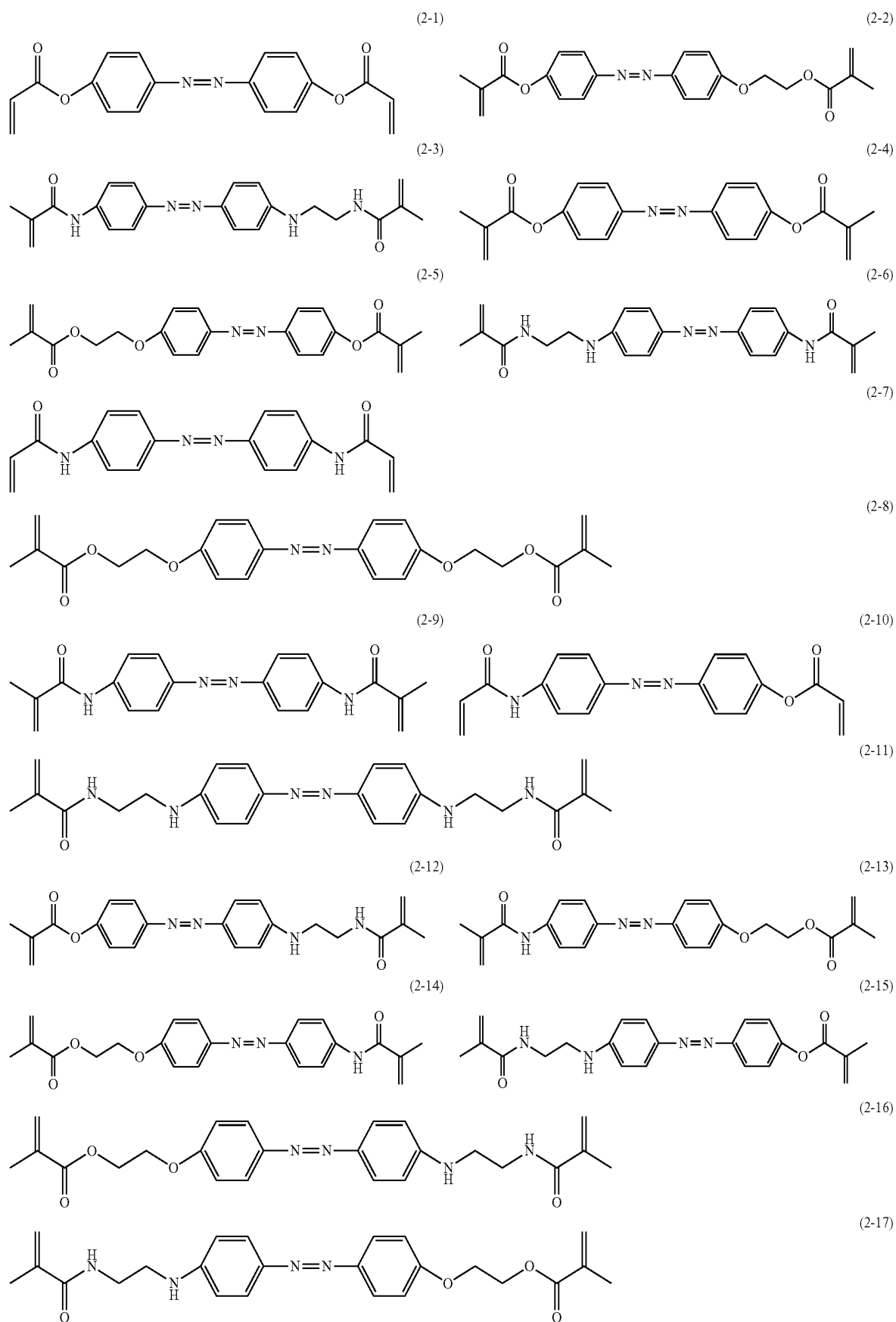
following formula (1) (hereinafter, also referred to as the monomer (1)). As described above, a polymer of the monomer (1) is likely to separate from the liquid crystal layer 40, which means that use of the monomer (1) enables easy formation of the polymer layers 31 and 51, reducing the ultraviolet light irradiation dose for the liquid crystal layer 40 in the polymer layer formation. This also contributes to maintenance of the VHR and reduction of residual DC voltage generation in long-term use. Also, since polarized ultraviolet light application imparts alignability to the monomer (1), use of the monomer (1) and application of polarized ultraviolet light (this process requires heating at the T_{ni} of the liquid crystal material or higher) enable formation of the polymer layers 31 and 51 capable of achieving the horizontal alignment. The liquid crystal display device of the present embodiment may therefore be in the vertical alignment mode, but is preferably in the horizontal alignment mode. Each of the polymer layers 31 and 51 preferably align liquid crystal compounds in the liquid crystal layer 40 in the direction parallel to the surfaces of the substrates 20 and 60. Aligning liquid crystal compounds in the liquid crystal layer 40 in the direction parallel to the surfaces of the substrates 20 and 60 means that the pre-tilt angle of the liquid crystal material from the surfaces of the substrates 20 and 60 is 0° or greater and 5° or smaller, preferably 0° or greater and 3° or smaller. The second polymer may include monomer (1)-derived units as main units or may be composed of the monomer (1)-derived units alone. At least some molecules of the second polymer in each of the polymer layers 31 and 51 are covalently bonded through a reaction to the functional group represented by the formula (A-1) (and radicals thereof) or the functional group represented by the formula (A-2) (and radicals thereof).



[0088] In the formula, P¹ and P² are the same as or different from each other, and each represent an acryloyloxy, methacryloyloxy, acryloylamino, methacryloylamino, vinyl, or vinyloxy group, Sp¹ and Sp² are the same as or different from each other, and each represent a C1-C10 linear, branched, or cyclic alkylene or alkenylene group, or a direct bond, L¹ and L² are the same as or different from each other, and each represent a —NH— group, a —N(CH₃)— group, a —O— group, a —S— group, or a direct bond, and at least one hydrogen atom in each phenylene group is optionally replaced.

[0089] In the formula (1), at least one hydrogen atom of each phenylene group is optionally replaced by a halogen atom (preferably a fluorine or chlorine atom), or a methyl, methoxy, ethyl, or ethoxy group.

[0090] Specific preferred examples of the monomer (1) include monomers represented by any of the following formulas (2-1) to (2-17). These may be used alone or in combination with each other.



[0091] The liquid crystal layer **40** contains a liquid crystal material (nematic liquid crystal) containing at least one liquid crystal compound (liquid crystal molecules) and exhibiting a nematic phase. The liquid crystal material transforms from the nematic phase into an isotropic phase when the temperature being increased from the nematic temperature reaches a certain critical temperature (nematic-isotropic phase transition temperature (T_{ni})). The nematic-isotropic phase transition temperature of the liquid crystal material may be any temperature, such as a temperature of 70° C. to 110° C. The T_{ni} is that of the liquid crystal material before the above monomer is added. The liquid crystal layer **40** contains liquid crystal compounds aligned in a predetermined direction with no voltage applied. The predetermined direction (initial alignment direction) is appropriately determined according to the liquid crystal driving mode. For example, in the horizontal alignment mode, the liquid crystal compounds are aligned in a direction substantially parallel to a main surface of each of the substrates **20** and **60** with no voltage applied. In the vertical alignment mode, the liquid crystal compounds are aligned in a direction substantially perpendicular to the main surface of each of the substrates **20** and **60** with no voltage applied.

[0092] The anisotropy of dielectric constant ($\Delta\epsilon$) represented by the following formula of the liquid crystal material and the liquid crystal compound may be positive or negative. The liquid crystal material may also contain a liquid crystal compound having no polarity, i.e., having an anisotropy $\Delta\epsilon$ of substantially 0 (neutral liquid crystal compound). Examples of the neutral liquid crystal compound include liquid crystal compounds having an alkene structure.

$$\Delta\epsilon = (\text{dielectric constant in major axis direction}) - (\text{dielectric constant in minor axis direction})$$

[0093] The thin-film transistor (TFT) substrate **60** can be an active matrix substrate usually used in the field of liquid crystal display panels. The liquid crystal driving mode for the liquid crystal display device of the present embodiment may be any mode such as the twisted nematic (TN) mode, the electrically controlled birefringence (ECB) mode, a horizontal alignment mode including the FFS mode and the IPS mode, or a vertical alignment mode including the 4D-RTN mode and the multi-domain vertical alignment (MVA) mode.

[0094] In the case where the liquid crystal driving mode of the liquid crystal display device of the present embodiment is the FFS mode, the TFT substrate **60** includes, for 1.0 example, a supporting substrate, a common electrode (planar electrode) disposed on the liquid crystal layer **40** side of the supporting substrate, an insulating film covering the common electrode, and pixel electrodes (comb electrodes) disposed on the liquid crystal layer **40** side of the insulating film. This structure can generate horizontal electric fields (fringe electric fields) in the liquid crystal layer **40** by applying voltage between the common electrode and the pixel electrodes constituting a pair of electrodes. Thus, controlling the voltage applied between the common electrode and the pixel electrodes enables control of the alignment of the liquid crystals in the liquid crystal layer **40**.

[0095] In the case where the liquid crystal driving mode for the liquid crystal display device of the present embodiment is the IPS mode, the liquid crystal display device applies voltage to the pair of comb electrodes disposed in the TFT substrate **60** to generate horizontal electric fields in the

liquid crystal layer **40**, thereby controlling the alignment of the liquid crystals in the liquid crystal layer **40**.

[0096] In the case where the liquid crystal driving mode for the liquid crystal display device of the present embodiment is the vertical alignment mode, the TFT substrate **60** includes pixel electrodes, and the counter substrate **20** includes a common electrode. The liquid crystal display device applies voltage between the common electrode and the pixel electrodes to generate vertical electric fields in the liquid crystal layer **40**, thereby controlling the alignment of the liquid crystals in the liquid crystal layer **40**. In the 4D-RTN mode, the alignment films **30** and **50** are subjected to alignment treatment in opposite (antiparallel) directions in each pixel, and the TFT substrate **60** and the counter substrate **20** are bonded to each other such that the alignment treatment directions for the alignment films **30** and **50** are perpendicular to each other. This can define four alignment directions (domains), which are different from each other, in each pixel. In the MVA mode, alignment control structures such as ribs and slits are provided to the TFT substrate **60** and the counter substrate **20** to define different alignment directions (domains) in each pixel.

[0097] The second linear polarizer **70** can be the same polarizer as that used for the first linear polarizer **10**. The transmission axis of the first linear polarizer **10** and the transmission axis of the second linear polarizer **70** are preferably perpendicular to each other. This configuration sets the first linear polarizer **10** and the second linear polarizer **70** in crossed Nicols, achieving a favorable black display state with no voltage applied.

[0098] The backlight **80** may be of any type and may be, for example, an edge-lit backlight or a direct-lit backlight. The backlight **80** may utilize any light source such as a light emitting diode (LED) or a cold cathode fluorescent lamp (CCFL). The amount of light emitted from the backlight **80** and transmitted by the light crystal panel is controlled by the voltage applied to the liquid crystal layer **40**.

[0099] The liquid crystal display device of Embodiment 1 may include any other members. For example, with an anti-reflection film disposed on the viewing surface side of the first linear polarizer **10**, the reflectance of the liquid crystal panel can be further decreased. The anti-reflection film is preferably a moth-eye film having a surface structure like a moth's eye.

[0100] The method for manufacturing the liquid crystal display device of the present embodiment is described.

[0101] First, the counter substrate **20** and the TFT substrate **60** are produced by common methods to prepare the counter substrate **20** and the TFT substrate **60** as a pair of substrates (preparation process).

[0102] The preparation is followed by applying an alignment agent containing a first polymer (polyimide-based first polymer) containing in its main chain at least one selected from a polyamic acid structure and a polyimide structure to the surface of each of the substrates **20** and **60** so as to form an alignment film (film forming process). Specifically, an alignment film agent is prepared by dissolving a polyimide-based first polymer in a solvent (for example, organic solvent). The polyimide-based first polymer preferably contains at least one functional group represented by any of the formulas (3-1) to (3-7). To the surface of each of the substrates **20** and **60** is applied the alignment agent by a method such as roll coating, spin coating, printing, or ink jetting. The surface of each of the substrates **20** and **60** is

heated to volatilize the solvent in the alignment agent, whereby the alignment films **30** and **50** are formed. The heating may include two processes of pre-baking and post-baking. The post-baking may include two processes, so that a total of three heating processes may be performed. In the case of using the polyimide-based first polymer, the post-baking may involve partial imidization of a polyamic acid structure into a polyimide structure.

[0103] Thereafter, the alignment films **30** and **50** may or may not be subjected to alignment treatment. Examples of the alignment treatment include rubbing and photoalignment treatments such as ultraviolet light irradiation. In a typical horizontal alignment mode, the horizontal alignment films are subjected to an alignment treatment. In the present embodiment employing the monomer (1), the alignment treatment in this process may be omitted. Even without the alignment treatment, irradiating the liquid crystal layer **40** with polarized ultraviolet light in the later-described polymer layer formation enables formation of the polymer layers **31** and **51** capable of achieving the horizontal alignment.

[0104] Between the substrates **20** and **60** having formed thereon the alignment films **30** and **50**, respectively, is formed the liquid crystal layer **40** by sealing a liquid crystal composition containing a liquid crystal material and at least one monomer (hereinafter, also referred to as an additional monomer) including at least one monomer represented by the formula (1) (monomer (1)) (liquid crystal layer formation). The liquid crystal layer formation is achieved by vacuum filling or one drop filling. The amount of the additional monomer to be introduced is preferably 1 to 30 wt %, more preferably 5 to 25 wt %, relative to the whole liquid crystal composition. The additional monomer may include the monomer (1) as a main monomer or may be composed of the monomer (1) alone. In the case of employing the vacuum filling, processes are performed in the following order: application of a sealant; bonding of the substrates **20** and **60**; curing of the sealant; injection of the liquid crystal composition; and sealing of the injection ports. In the case of employing the one drop filling, processes are performed in the following order: application of a sealant; dropping of the liquid crystal composition; bonding of the substrates **20** and **60**; and curing of the sealant. These result in a liquid crystal cell filled with the liquid crystal material.

[0105] The liquid crystal layer **40** between the substrates **20** and **60** is then heated to transform the liquid crystal material into an isotropic phase (isotropic phase treatment). The heating temperature here may be any temperature higher than the nematic-isotropic phase transition temperature of the liquid crystal material and is, for example, 80° C. to 150° C. The heating duration is, for example, 30 to 60 minutes. The isotropic phase treatment is followed by cooling the liquid crystal cell to room temperature.

[0106] After the isotropic phase treatment, the liquid crystal layer **40** is irradiated with ultraviolet light to form the polymer layers **31** and **51** from the second polymer obtained by polymerizing the additional monomer including the monomer (1) between the liquid crystal layer **40** and the alignment films **30** and **50** (polymer layer formation), respectively. Also, in the polymer layer formation, the second polymer is covalently bonded to the polyimide-based first polymer. The ultraviolet light applied is preferably polarized ultraviolet light, more preferably linearly polarized ultraviolet light, in the case of the horizontal alignment

mode, and is preferably unpolarized ultraviolet light in the case of the vertical alignment mode.

[0107] The ultraviolet light may have a wavelength of 200 to 430 nm. The lower limit of the wavelength is more preferably 250 nm, and the upper limit is more preferably 380 nm. The ultraviolet light may be applied with an irradiation dose of 0.3 to 20 J/cm². The lower limit of the irradiation dose is more preferably 1 J/cm, and the upper limit is more preferably 5 J/cm².

[0108] In the case of the horizontal alignment mode, the polymer layer formation preferably includes applying polarized ultraviolet light while heating the liquid crystal layer **40** at a temperature equal to or higher than the T_{ni} of the liquid crystal material. This can prevent the liquid crystal material in the liquid crystal layer **40** from changing the state of the applied polarized ultraviolet light, successfully achieving the horizontal alignment of the liquid crystal compounds. The lower limit of the heating temperature is T_{ni}+3° C. or higher. The upper limit of the heating temperature is preferably T_{ni}+20° C. or lower for minimization of heat deterioration of the liquid crystal material.

[0109] In the case of the vertical alignment mode, the polymer layer formation may include applying unpolarized ultraviolet rays while applying voltage (for example, 4 to 10 V) to the liquid crystal layer **40**.

[0110] Performing the polymer layer formation after the liquid crystal layer formation enables bonding of the substrates **20** and **60**, holding the liquid crystal layer **40** in between, to each other with the sealant and formation of the polymer layers **31** and **51** in a region surrounded by the sealant in a plan view.

[0111] The above processes are followed by bonding of polarizers and mounting of members such as a controller, a power source, and a backlight. Thereby, the liquid crystal display device of the present embodiment is completed.

[0112] An embodiment of the present invention was described above. Each and every feature of the embodiment is applicable to all the aspects of the present invention.

[0113] The present invention is described in more detail below based on examples and comparative examples. The examples, however, are not intended to limit the scope of the present invention.

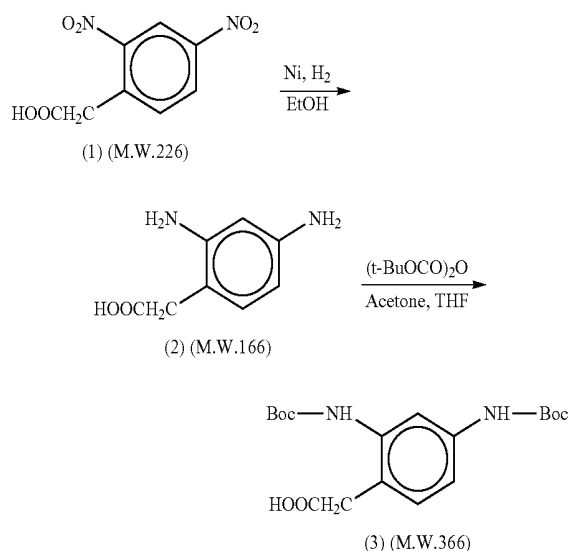
Synthesis Example 1

[0114] The following shows an exemplary synthesis of a diamine monomer containing a 2-hydroxy-2-methyl-1-phenyl-propan-1-one functional group in a side chain.

Process A

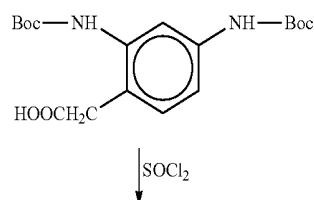
[0115] An amount of 3 g of dinitrophenyl acetate (13.3 mmol, compound (1)) was dissolved in 8 mL of SOLMIX AP-I, followed by addition of 0.06 g of Raney Ni to the solution. The mixture was fed into an autoclave. The system was purged with hydrogen and left to stand overnight at room temperature under a pressure of 0.4 MPa. High performance liquid chromatography (HPLC) was used to confirm that the reaction was completed, and then the reaction liquid was filtered through Celite. The filtrate was concentrated until no filtrate was observed. The thus-obtained crude liquid was distilled under reduced pressure, so that 1.98 g of 2,4-diaminophenyl acetate (2) (11.9 mmol, yield: 90%) was obtained. Then, 1.8 g of the compound (2) (10.8 mmol) was dissolved in 5 mL of acetone, followed by dropwise addition

of t-butoxycarbonyl anhydride (5 g/THF 5 mL) into the solution. The resulting mixture was heated to the reflux temperature and left to stand overnight. After the completion of the reaction, the reaction liquid was concentrated and dried, whereby a Boc compound (3) (3.73 g, 10.2 mmol, yield: 94%) was obtained.

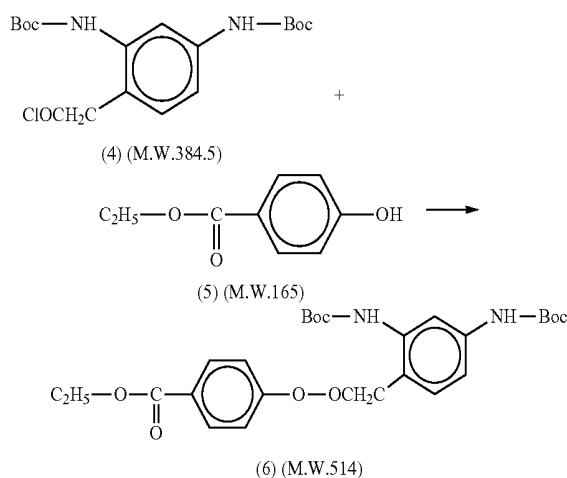


Process B

[0116] Into a benzene solution (30 mL) containing 3.5 g (9.56 mmol) of the Boc compound (3) was dropwise added thionyl chloride, so that an acid chloride compound represented by the following formula (4) (3.42 g, 8.89 mmol, yield: 93%) was synthesized. Into a benzene solution (30 mL) containing 1.65 g (10 mmol) of ethyl 4-hydroxybenzoate represented by the following formula (5) and 1.5 g (15 mmol) of triethylamine was dropwise added a benzene solution (20 mL) containing 3.3 g (8.5 mmol) of the acid chloride compound represented by the following formula (4) at room temperature in a nitrogen atmosphere. The mixture was reacted for two hours at room temperature. After completion of the reaction, impurities were extracted with water, and the residue was purified by column chromatography (toluene/ethyl acetate (4/1)), whereby 4.11 g of the target compound represented by the following formula (6) was obtained (yield: 80%).

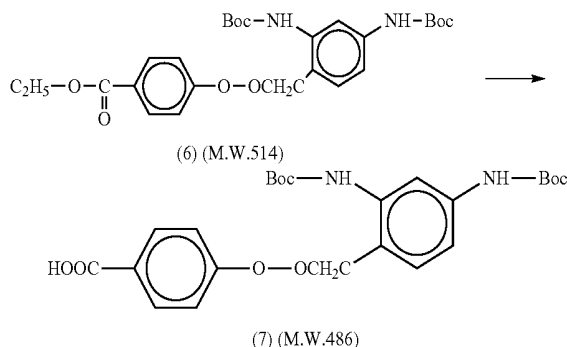


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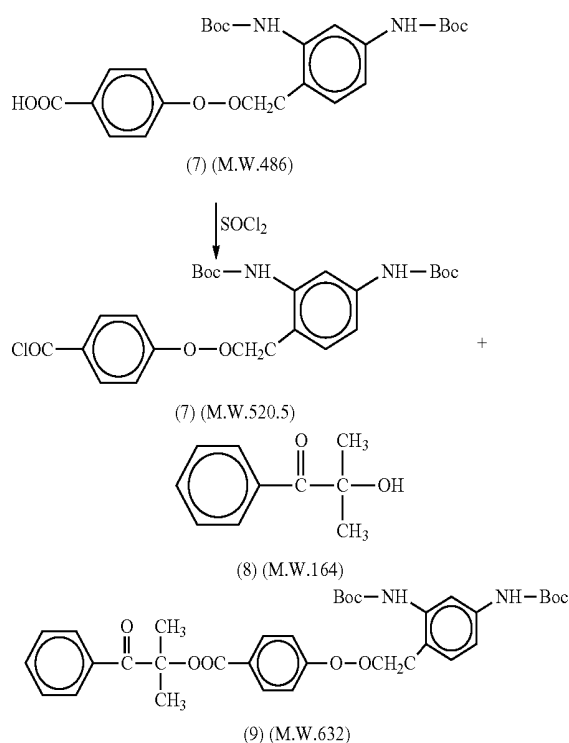
Process C

[0117] Into a THF/methanol mixed solution (20 mL) containing 2.57 g (5 mmol) of a compound represented by the formula (6) was dropwise added a sodium hydroxide aqueous solution and then hydrochloric acid. The resulting mixture was stirred for one hour, so that a carboxylic acid compound represented by the following formula (7) was synthesized (2.33 g, 4.8 mmol).



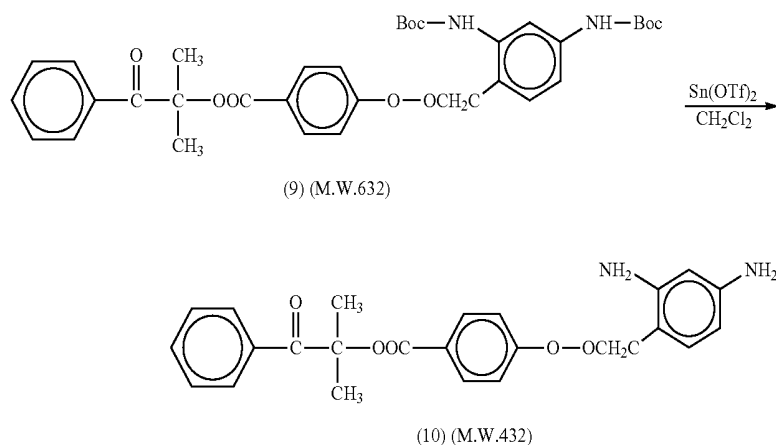
Process D

[0118] Thionyl chloride was dropwise added into a benzene solution (30 mL) containing 2.19 g (4.5 mmol) of a compound represented by the formula (7), so that an acid chloride compound (2.24 g, 4.3 mmol, yield: 96%) was synthesized. Into a benzene solution (30 mL) containing 0.82 g (5 mmol) of 2-hydroxy-2-methyl-1-phenyl-propan-1-one represented by the following formula (8) and 1.5 g (15 mmol) of triethylamine was dropwise added a benzene solution (20 mL) containing 2.24 g (4.3 mmol) of an acid chloride compound represented by the following formula (7) at room temperature in a nitrogen atmosphere. The mixture was reacted for two hours at room temperature. After completion of the reaction, impurities were extracted with water, and the residue was purified by column chromatography (toluene/ethyl acetate (4/1)), whereby 2.26 g of the target compound represented by the following formula (9) (3.57 mmol, yield: 83%) was obtained.

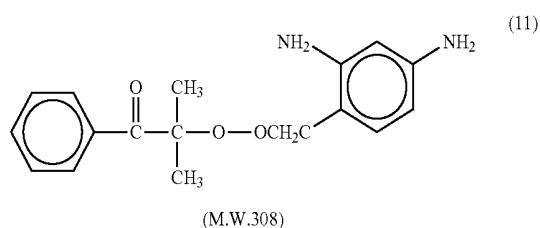


Process E

[0119] The compound (9) was dissolved in methylene chloride, followed by adding tin(II) trifluoromethanesulfonate ($\text{Sn}(\text{OTf})_2$) in portions to the solution at 0°C . The mixture was reacted at room temperature, and then neutralized with 5% NaHCO_3 aq. The mixture was washed with water until it had a neutral pH. The organic layer was dried over anhydrous magnesium sulfate and filtered through Celite. The filtrate was concentrated, whereby the target diamine monomer (10) (1.13 g, 2.61 mmol, yield: 73%) was obtained.



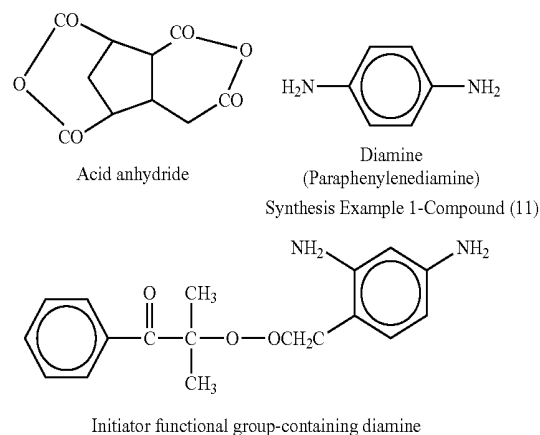
[0120] With the compound (4) and the compound (8) in place of the compound (7), a diamine monomer represented by the following formula (11) can be synthesized in the same manner.



Condensation Polymerization Example 1

[0121] The following shows an exemplary synthesis with 10 mol % of a polymerization initiator group-containing diamine unit introduced.

[0122] The following acid anhydride (0.10 mol) was added to a γ -butyrolactone solution containing the following paraphenylenediamine (0.09 mol) and the following 2-hydroxy-2-methyl-1-phenylpropan-1-one functional group-containing diamine (0.01 mol), and the mixture was reacted at 60°C . for 12 hours, whereby a polyamic acid having a random structure was obtained.

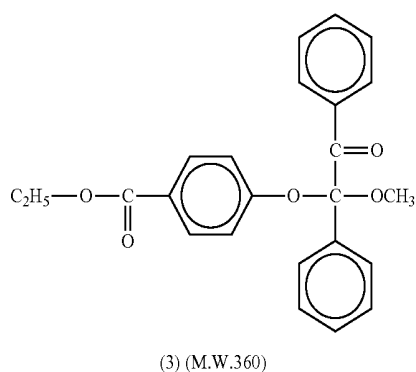
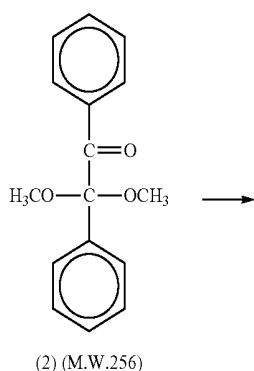
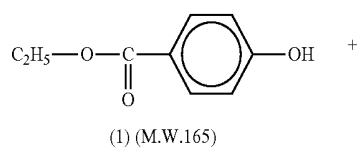


Synthesis Example 2

[0123] The following shows an exemplary synthesis of a diamine monomer containing a benzyl ketal-based initiator in a side chain.

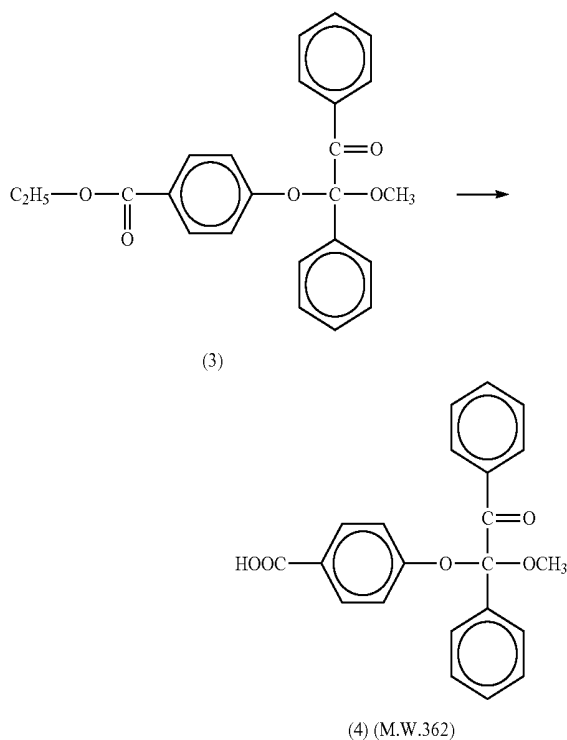
Process A

[0124] A benzene solution (5 mL) containing 0.64 g (2.5 mmol) of benzyl dimethyl ketal represented by the following formula (2) was dropwise added into a benzene solution (20 mL) containing 0.42 g (2.5 mmol) of ethyl 4-hydroxybenzoate (the following formula (1)) and 0.5 g (5 mmol) of triethylamine at room temperature in a nitrogen atmosphere. The mixture was reacted at room temperature for two hours. After completion of the reaction, the impurities were extracted with water, and the residue was purified by column chromatography (toluene/ethyl acetate (4/1)), whereby 0.878 g (yield: 90%) of the target compound represented by the following formula (3) was obtained.

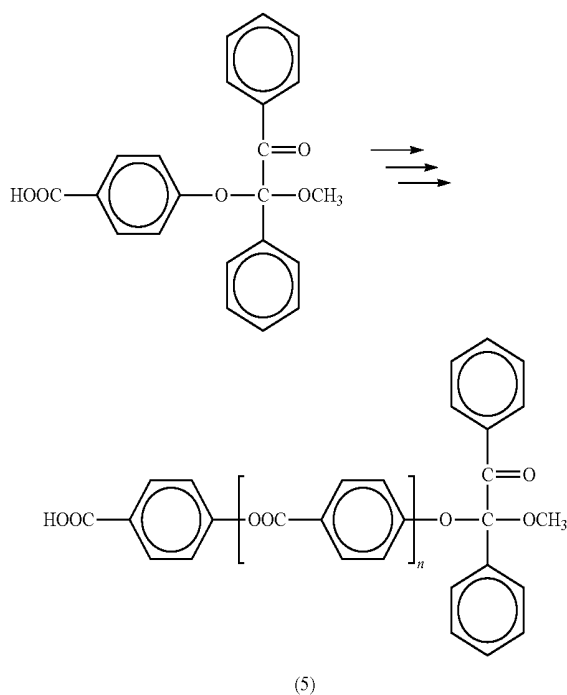


Process B

[0125] Into a THF/methanol mixed solution (20 mL) containing 0.78 g (2 mmol) of the compound represented by the formula (3) was dropwise added a sodium hydroxide aqueous solution and then hydrochloric acid. The mixture was stirred for one hour, so that a carboxylic acid compound represented by the following formula (4) was synthesized (0.69 g, 1.9 mmol).



[0126] A compound represented by the following formula (5) was synthesized by repeating the processes A and B.

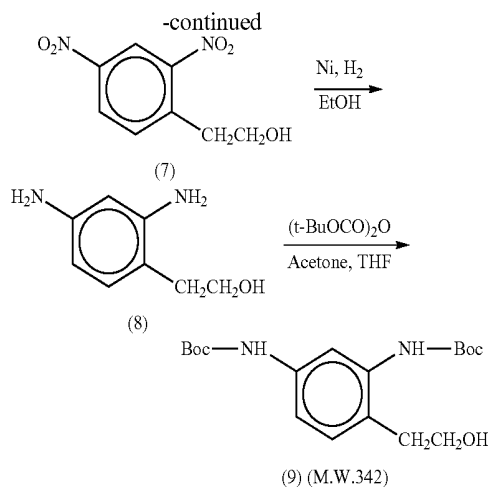
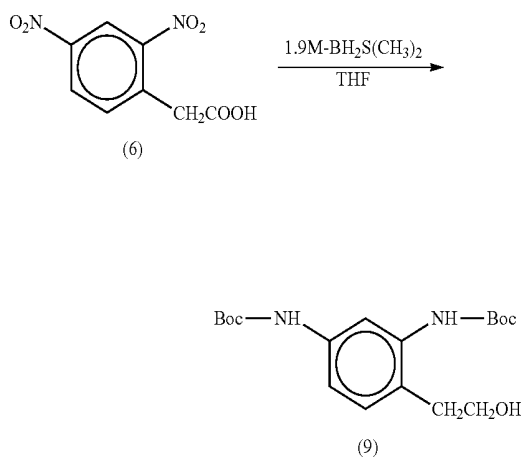


$n = 0, 1, 2, 3$

Process C

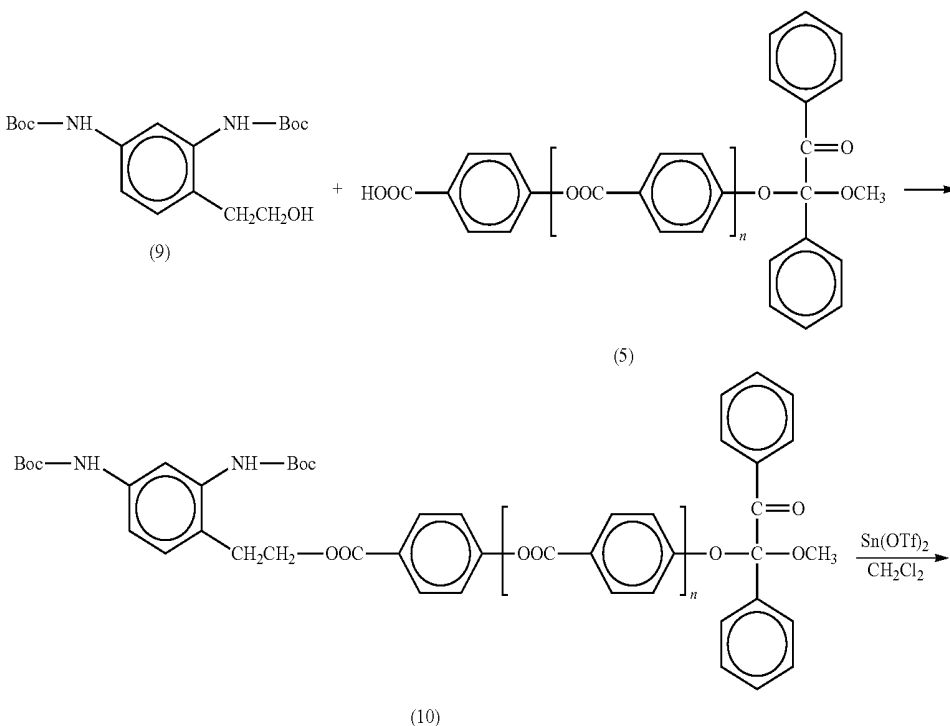
[0127] An amount of 3 g of dinitrophenyl acetate (compound (6)) was dissolved in 20 mL of THF, followed by

dropwise addition of borane dimethyl sulfide-toluene solution (7 mL) into the solution. The resulting mixture was left to stand overnight at room temperature. The reaction was terminated by dropwise adding 10 mL of a 50% methanol aqueous solution to the mixture. The target component was extracted with 10 mL of chloroform, washed with 5% sodium bicarbonate water and water, and concentrated until dissolution into the organic layer stopped. The resulting liquid was dissolved in 20 mL of chloroform, and the solution was purified by aluminum column chromatography. The distillate was concentrated. A toluene/n-heptane solution (6/4) was added to the concentrate, and a component was heat-extracted at 70° C. and isolated. Decantation was used to obtain the upper layer of the component. The component was cooled to obtain 2,4-dinitrophenyl ethanol (7) (1.2 g, yield: 42.7%). An amount of 0.4 g of the compound (7) was dissolved in 8 mL of SOLMIX AP-T, followed by addition of 0.06 g of Raney Ni to the solution. The mixture was fed into an autoclave. The system was purged with hydrogen and left to stand overnight at room temperature under a pressure of 0.4 MPa. HPLC was used to confirm that the reaction was completed, and then the reaction liquid was filtered through Celite. The filtrate was concentrated until no filtrate was observed. The thus-obtained crude liquid was distilled under reduced pressure, so that 0.69 g (yield: 80%) of 2,4-diaminophenyl ethanol (8) was obtained. An amount of 0.6 g of the compound (8) was dissolved in 5 mL of acetone, followed by dropwise addition of a t-butoxycarbonyl anhydride (1.8 g/THF 5 mL). The resulting mixture was heated to the reflux temperature and left to stand overnight. After the completion of the reaction, the reaction liquid was concentrated and dried, whereby a Boc compound (9) (0.13 g, yield: 94%) was obtained.

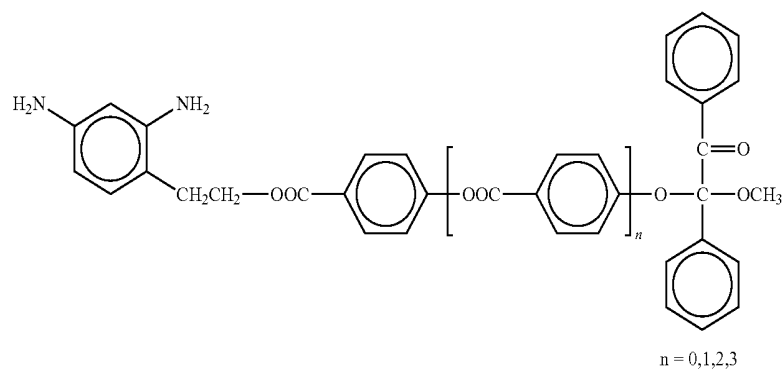


Process D

[0128] The Boc compound (9) and a carboxylic acid compound represented by the following formula (5) were reacted by the same process as Process A, so that a compound represented by the following formula (10) was synthesized. The Boc compound was then converted back to a diamine, so that the target monomer (11) was synthesized. The following shows how to synthesize the monomer (11) from the compound (10). The compound (10) was dissolved in methylene chloride, followed by adding tin(II) trifluoromethanesulfonate (Sn(OTf)₂) in portions to the solution at 0° C. The mixture was reacted at room temperature, and then neutralized with 5% NaHCO₃ aq. The mixture was washed with water until it had a neutral pH. The organic layer was dried over anhydrous magnesium sulfate and filtered through Celite. The filtrate was concentrated, whereby a diamine monomer (11) was obtained.



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Condensation Polymerization Example 2

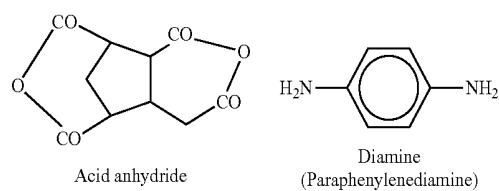
[0129] The following shows an exemplary synthesis with 10 mol % of a benzyl ketal-based polymerization initiator group-containing diamine unit introduced.

[0130] The following acid anhydride (0.10 mol) was added to a γ -butyrolactone solution containing the following paraphenylenediamine (0.09 mol) and the following benzyl ketal-based polymerization initiator group-containing diamine (0.01 mol), and the mixture was reacted at 60° C. for 12 hours, whereby a polyamic acid having a random structure was obtained.

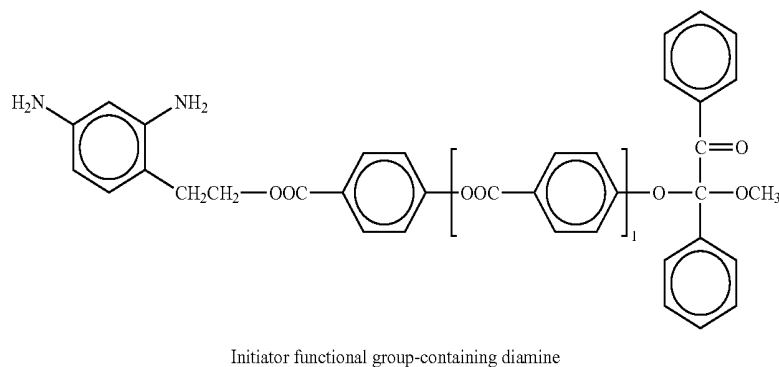
Condensation Polymerization Example 3

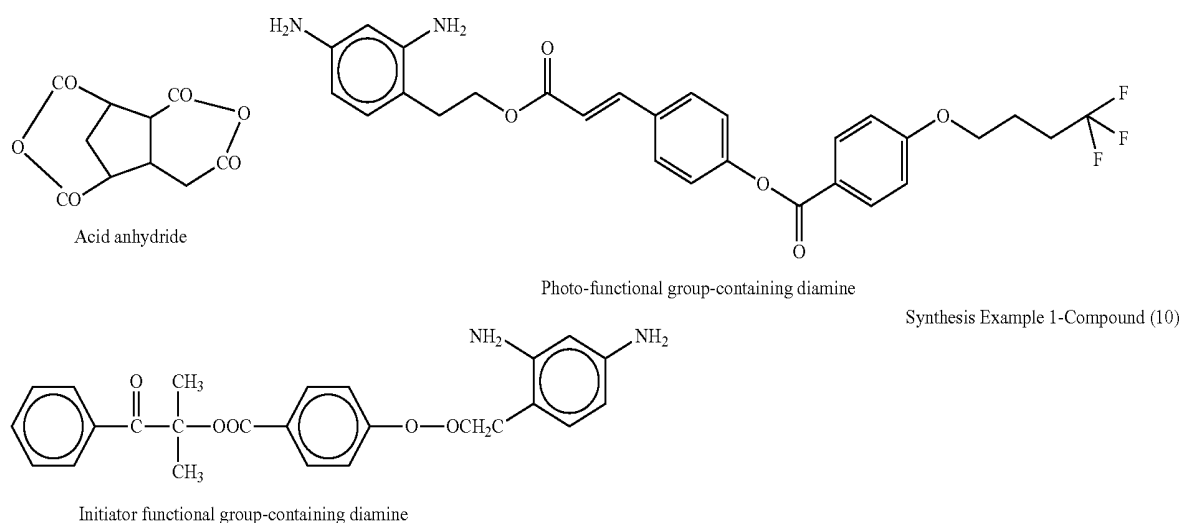
[0131] The following shows an exemplary synthesis with 10 mol % of a polymerization initiator group-containing diamine unit introduced.

[0132] The following acid anhydride (0.10 mol) was added to a γ -butyrolactone solution containing the following photo-functional group-containing diamine (0.09 mol) and the following 2-hydroxy-2-methyl-1-phenyl-propan-1-one functional group-containing diamine (0.01 mol), and the mixture was reacted at 60° C. for 12 hours, whereby a polyamic acid having a random structure was obtained.



Synthesis Example 2-Compound (11)



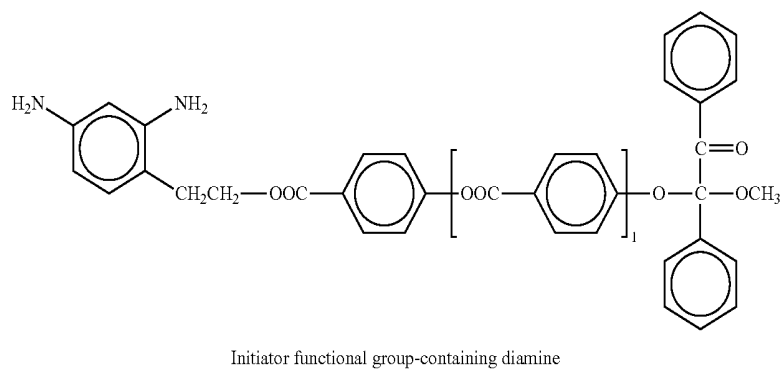
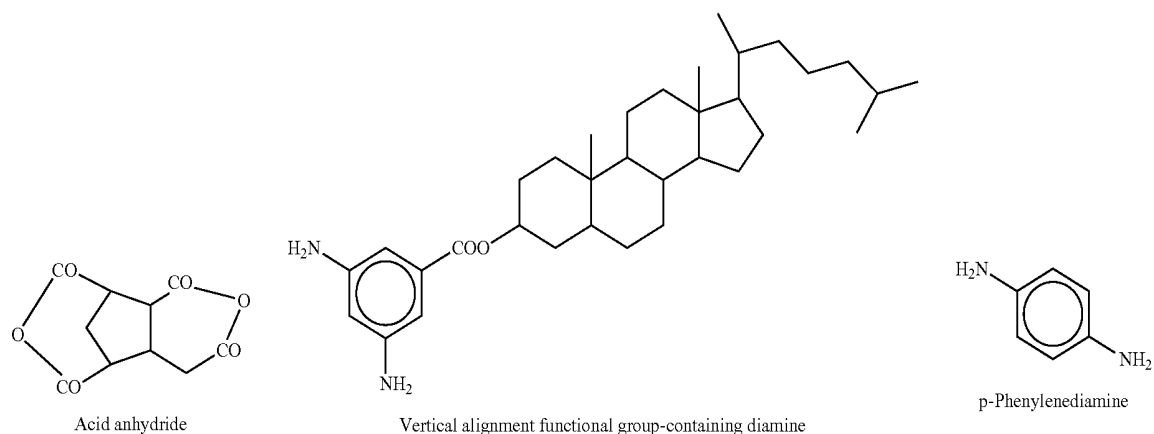


Condensation Polymerization Example 4

[0133] The following shows an exemplary synthesis with 10 mol % of a polymerization initiator group-containing diamine unit introduced.

[0134] The following acid anhydride (0.10 mol) was added to a γ -butyrolactone solution containing the following

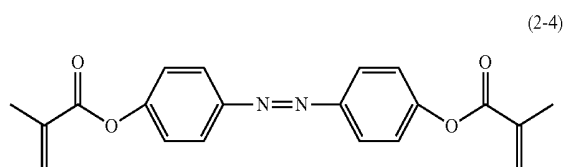
vertical alignment functional group-containing diamine (0.03 mol), the following p-phenylene diamine (0.06 mol), and the following polymerization initiator group-containing diamine (0.01 mol), and the mixture was reacted at 60° C. for 12 hours, whereby a polyamic acid having a random structure was obtained.



Example 1-1

(Production of Liquid Crystal Cell)

[0135] A TFT substrate and a counter substrate with no electrode for the FFS mode were prepared. A horizontal alignment film was formed on each of the TFT substrate and the counter substrate using a polyamic acid containing in a side chain an initiator functional group synthesized as in Condensation Polymerization Example 1. The amount *m* of the initiator functional group-containing diamine unit introduced into the polyamic acid was 20 mol %. A seal was applied to one of the substrates, and beads were scattered on the counter substrate. The substrates were then bonded to each other. Between the substrates was injected a liquid crystal material having positive anisotropy of dielectric constant ($T_{ni}=85^{\circ}\text{C.}$, $\Delta n=0.11$, $\Delta\epsilon=6.5$). The liquid crystal material contained 0.6 wt % of an azobenzene group-containing bifunctional monomer represented by the following formula (2-4). The liquid crystal material injection was followed by heating at 130°C. and quenching. The liquid crystal cell was irradiated with linearly polarized ultraviolet rays having a wavelength of 365 nm for 30 minutes (irradiation dose: 3 J/cm^2) with a cell temperature of 90°C. to polymerize the monomer, and thereby an FFS mode liquid crystal cell was obtained.



Examples 1-2 to 1-5

[0136] FFS mode liquid crystal cells of Examples 1-2, 1-3, 1-4, and 1-5 were produced as in Example 1-1, except that the amount *m* of the initiator functional group-containing diamine unit introduced into the polyamic acid was changed to 40 mol %, 60 mol %, 80 mol %, and 100 mol %, respectively.

Comparative Example 1-1

[0137] An FFS mode liquid crystal cell of Comparative Example 1-1 was produced as in Example 1-1, except that the amount *m* of the initiator functional group-containing diamine unit introduced into the polyamic acid was changed to 0 mol %.

Comparative Example 1-2

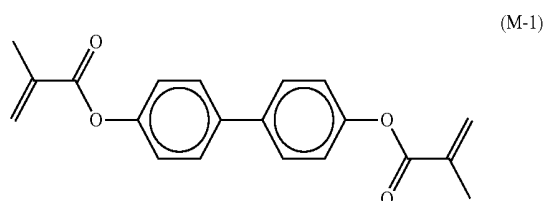
[0138] An FFS mode liquid crystal cell of Comparative Example 1-2 was produced as in Comparative Example 1-1, except that the cell temperature in the polarized ultraviolet light application for polymerization of the monomer was changed to room temperature (35°C.).

Comparative Examples 1-3 to 1-7

[0139] FFS mode liquid crystal cells of Comparative Examples 1-3, 1-4, 1-5, 1-6, and 1-7 were produced as in Examples 1-1, 1-2, 1-3, 1-4, and 1-5, respectively, except that the cell temperature in the polarized ultraviolet light application for polymerization of the monomer was changed to room temperature (35°C.).

Comparative Example 1-8

[0140] An FFS mode liquid crystal cell of Comparative Example 1-8 was produced as in Comparative Example 1-1, except that the liquid crystal material contained 0.6 wt % of a bifunctional monomer represented by the following formula (M-1) instead of the monomer represented by the formula (2-4).



Comparative Examples 1-9 to 1-13

[0141] FFS mode liquid crystal cells of Comparative Examples 1-9, 1-10, 1-1.1, 1-12, and 1-13 were produced as in Examples 1-1, 1-2, 1-3, 1-4, and 1-5, respectively, except that the liquid crystal material contained 0.6 wt % of a bifunctional monomer represented by the formula (M-1) instead of the monomer represented by the formula (2-4).

(Alignment Confirmation)

[0142] The alignment in each of the obtained liquid crystal cells was visually confirmed.

(Backlight Exposure Power-on Test)

[0143] Each of the produced liquid crystal cells was sandwiched between polarizers and then power was supplied to the workpiece on a backlight for 100 hours. The power supply conditions were 10 V and 30 Hz. The contrast ratio, the voltage holding ratio (VHR), and residual DC (rDC) voltage before and after the power supply to the workpiece on the backlight were measured. The contrast ratio was measured using BM-5A from Topcon Corporation (25°C.). The VHR was measured at 1 V (70°C.) using a VHR measurement system from Toyo Corp. The rDC voltage was measured at a DC offset voltage of 2 V by flicker elimination method.

[0144] The results are shown in the following Tables 1-1 to 1-3.

TABLE 1-1

Results of using azobenzene group-containing bifunctional monomer and applying polarized UV light at 90° C.							
		Comparative Example 1-1	Example 1-1	Example 1-2	Example 1-3	Example 1-4	Example 1-5
Initiator introduced (mol %)		m = 0	m = 20	m = 40	m = 60	m = 80	m = 100
Alignment state	Initial	Horizontally aligned	Horizontally aligned	Horizontally aligned	Horizontally aligned	Horizontally aligned	Horizontally aligned
Contrast ratio	Initial	820	820	820	820	820	820
	100 h	720	810	800	810	810	800
VHR (%)	Initial	99.2	99.2	99.3	99.3	99.3	99.1
	100 h	95.7	98.4	98.5	98.5	98.6	98.3
rDC voltage (mV)	Initial	20	20	20	20	30	40
	100 h	180	50	40	40	40	40

TABLE 1-2

Results of using azobenzene group-containing bifunctional monomer and applying polarized UV light at 35° C.							
		Comparative Example 1-2	Comparative Example 1-3	Comparative Example 1-4	Comparative Example 1-5	Comparative Example 1-6	Comparative Example 1-7
Initiator introduced (mol %)		m = 0	m = 20	m = 40	m = 60	m = 80	m = 100
Alignment state	Initial	Random	Random	Random	Random	Random	Random
Contrast ratio	Initial	(Unmeasurable due to unaligned state)					
VHR (%)	Initial	99.4	99.4	99.4	99.4	99.3	99.3
	100 h	97.7	99.3	99.3	99.3	99.1	98.9
rDC voltage (mV)	Initial	(Unmeasurable due to unaligned state)					

TABLE 1-3

Results of using biphenyl group-containing bifunctional monomer and applying polarized UV light at 90° C.							
		Comparative Example 1-8	Comparative Example 1-9	Comparative Example 1-10	Comparative Example 1-11	Comparative Example 1-12	Comparative Example 1-13
Initiator introduced (mol %)		m = 0	m = 20	m = 40	m = 60	m = 80	m = 100
Alignment state	Initial	Random	Random	Random	Random	Random	Random
Contrast ratio	Initial	(Unmeasurable due to unaligned state)					
VHR (%)	Initial	99.5	99.6	99.5	99.5	99.6	99.6
	100 h	98.8	99.5	99.5	99.5	99.3	99.4
rDC voltage (mV)	Initial	(Unmeasurable due to unaligned state)					

[0145] Only when the azobenzene group-containing bifunctional monomer represented by the formula (2-4) was used and polarized ultraviolet light was applied at 90° C. (equal to or higher than the T_{ni} of the liquid crystal material), the horizontal alignment was achieved. Also, Table 1-1 shows that in the cases where the polymerization initiator functional group was introduced into a side chain of a polyamic acid (partially imidized) constituting the alignment films (m=20 to 100), the backlight exposure turn-on test did not cause a contrast ratio decrease, a VHR decrease, or a rDC voltage increase.

[0146] Table 1-2 shows that the horizontal alignment was not achieved by applying polarized ultraviolet light at 35° C. (lower than the T_{ni} of the liquid crystal material) even when the azobenzene group-containing bifunctional monomer represented by the formula (2-4) was used. For achievement of the horizontal alignment mode, reducing scattering of polarized ultraviolet light, which is the light applied, by transforming the liquid crystal material into an isotropic phase seems to be important.

[0147] Also, Table 1-3 shows that the horizontal alignment was not achieved by applying polarized ultraviolet light at

90° C. when the biphenyl group-containing bifunctional monomer, which is a conventionally used monomer for PSA, was used (formula (M-1)). This is presumably because a biphenyl group was not alignable in the polymer layer formation using polarized ultraviolet light, and thus the liquid crystal alignability was not achieved.

Example 2-1

(Production of Liquid Crystal Cell)

[0148] A TFT substrate and a counter substrate with no electrode for the FFS mode were prepared. A horizontal alignment film was formed on each of the TFT substrate and the counter substrate using a polyamic acid containing in a side chain an initiator functional group synthesized as in Condensation Polymerization Example 2. The amount m of the initiator functional group-containing diamine unit introduced into the polyamic acid was 20 mol %. A seal was applied to one of the substrates, and beads were scattered on the counter substrate. The substrates were then bonded to each other. Between the substrates was injected a liquid crystal material having positive anisotropy of dielectric

constant ($T_{ni}=70^{\circ}\text{C.}$, $\Delta n=0.12$, $\Delta\epsilon=2.6$). The liquid crystal material contained 0.8 wt % of an azobenzene group-containing bifunctional monomer represented by the formula (2-4). The liquid crystal material injection was followed by heating at 130°C. and quenching. The liquid crystal cell was irradiated with linearly polarized ultraviolet rays having a wavelength of 365 nm for 30 minutes (irradiation dose: 3 J/cm^2) with a cell temperature of 80°C. to polymerize the monomer, and thereby an FFS mode liquid crystal cell was obtained.

Examples 2-2 to 2-5

[0149] FFS mode liquid crystal cells of Examples 2-2, 2-3, 2-4, and 2-5 were produced as in Example 2-1, except that the amount m of the initiator functional group-containing diamine unit introduced into the polyamic acid was changed to 40 mol %, 60 mol %, 80 mol %, and 100 mol %, respectively.

Comparative Example 2-1

[0150] An FFS mode liquid crystal cell of Comparative Example 2-1 was produced as in Example 2-1, except that the amount m of the initiator functional group-containing diamine unit introduced into the polyamic acid was changed to 0 mol %.

Comparative Example 2-2

[0151] An FFS mode liquid crystal cell of Comparative Example 2-2 was produced as in Comparative Example 2-1,

except that the cell temperature in the polarized ultraviolet light application for polymerization of the monomer was changed to room temperature (35°C.).

Comparative Examples 2-3 to 2-7

[0152] FFS mode liquid crystal cells of Comparative Examples 2-3, 2-4, 2-5, 2-6, and 2-7 were produced as in Examples 2-1, 2-2, 2-3, 2-4, and 2-5, respectively, except that the cell temperature in the polarized ultraviolet light application for polymerization of the monomer was changed to room temperature (35°C.).

Comparative Example 2-8

[0153] An FFS mode liquid crystal cell of Comparative Example 2-8 was produced as in Comparative Example 2-1, except that the liquid crystal material contained 0.8 wt % of a bifunctional monomer represented by the formula (M-1) instead of the monomer represented by the formula (2-4).

Comparative Examples 2-9 to 2-13

[0154] FFS mode liquid crystal cells of Comparative Examples 2-9, 2-10, 2-11, 2-12, and 2-13 were produced as in Examples 2-1, 2-2, 2-3, 2-4, and 2-5, respectively, except that the liquid crystal material contained 0.8 wt % of a bifunctional monomer represented by the formula (M-1) instead of the monomer represented by the formula (2-4).

[0155] The liquid crystal cells were subjected to the same evaluations as in Example 1-1 and the other examples. The results are shown in the following Tables 2-1 to 2-3.

TABLE 2-1

Results of using azobenzene group-containing bifunctional monomer and applying polarized UV light at 80°C.							
Comparative							
	Example 2-1	Example 2-1	Example 2-2	Example 2-3	Example 2-4	Example 2-5	
Initiator introduced (mol %)	$m = 0$	$m = 20$	$m = 40$	$m = 60$	$m = 80$	$m = 100$	
Alignment state	Initial	Horizontally aligned	Horizontally aligned	Horizontally aligned	Horizontally aligned	Horizontally aligned	Horizontally aligned
Contrast ratio	Initial	820	840	850	850	830	810
	100 h	750	820	820	820	820	800
VHR (%)	Initial	99.2	99.2	99.1	99.2	99.1	98.6
	100 h	95.2	98.6	98.8	98.8	98.8	98.2
rDC voltage (mV)	Initial	10	10	10	20	20	20
	100 h	190	40	30	30	30	20

TABLE 2-2

Results of using azobenzene group-containing bifunctional monomer and applying polarized UV light at 35°C.						
	Comparative Example 2-2	Comparative Example 2-3	Comparative Example 2-4	Comparative Example 2-5	Comparative Example 2-6	Comparative Example 2-7
Initiator introduced (mol %)	$m = 0$	$m = 20$	$m = 40$	$m = 60$	$m = 80$	$m = 100$
Alignment state	Initial	Random	Random	Random	Random	Random
Contrast ratio	Initial	(Unmeasurable due to unaligned state)				
VHR (%)	Initial	99.4	99.3	99.23	99.3	99.2
	100 h	97.5	98.9	99.0	98.9	98.4
rDC voltage (mV)	Initial	(Unmeasurable due to unaligned state)				

TABLE 2-3

Results of using biphenyl group-containing bifunctional monomer and applying polarized UV light at 80° C.						
	Comparative Example 2-8	Comparative Example 2-9	Comparative Example 2-10	Comparative Example 2-11	Comparative Example 2-12	Comparative Example 2-13
Initiator introduced (mol %)	m = 0	m = 20	m = 40	m = 60	m = 80	m = 100
Alignment state	Initial Random	Initial Random	Initial Random	Initial Random	Initial Random	Initial Random
Contrast ratio	Initial		(Unmeasurable due to unaligned state)			
VHR (%)	Initial	99.1	99.1	98.8	99.0	99.1
	100 h	96.2	98.7	98.6	98.7	98.6
rDC voltage (mV)	Initial		(Unmeasurable due to unaligned state)			

[0156] Only when the azobenzene group-containing bifunctional monomer represented by the formula (2-4) was used and polarized ultraviolet light was applied at 80° C. (equal to or higher than the T_{ni} of the liquid crystal material), the horizontal alignment was achieved. Also, Table 2-1 shows that in the cases where a polymerization initiator functional group was introduced into a side chain of a polyamic acid (partially imidized) constituting the alignment film (m=20 to 100), the backlight exposure turn-on test did not cause a contrast ratio decrease, a VHR decrease, or a rDC voltage increase.

[0157] Table 2-2 shows that the horizontal alignment was not achieved by applying polarized ultraviolet light at 35° C. (lower than the T_{ni} of the liquid crystal material) even when the azobenzene group-containing bifunctional monomer represented by the formula (2-4) was used. For achievement of the horizontal alignment mode, reducing scattering of polarized ultraviolet light, which is the light applied, by transforming the liquid crystal material into an isotropic phase seems to be important.

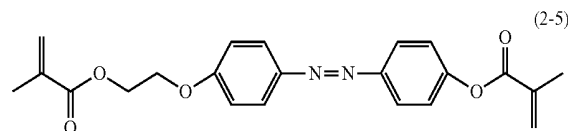
[0158] Also, Table 2-3 shows that the horizontal alignment was not achieved by applying polarized ultraviolet light at 80° C. when the biphenyl group-containing bifunctional monomer, which is a conventionally used monomer for PSA, was used (formula (M-1)). This is presumably because a biphenyl group was not alignable in the polymer layer formation using polarized ultraviolet light, and thus the liquid crystal alignability was not achieved.

Example 3-1

(Production of Liquid Crystal Cell)

[0159] A pair of substrates each having an ITO electrode on its entire surface was prepared. A vertical photo-alignment film was formed on each of the substrates using a polyamic acid containing in a side chain an initiator functional group synthesized as in Condensation Polymerization Example 3 and a vertical photo-functional group. The amount m of the initiator functional group-containing diamine unit introduced into the polyamic acid was 10 mol %. The film formation includes pre-baking the applied alignment agent at 80° C. for five minutes and then post-baking it at 200° C. for 40 minutes. The alignment agent was then irradiated with polarized ultraviolet light with an intensity of 30 mJ/cm² for an alignment treatment which enables the alignment films to give a pre-tilt angle of 88.5° (substantially vertical alignment). A seal was applied to one of the substrates, and beads were scattered on the other substrate. The substrates were then bonded to each other. Between the substrates was injected a liquid crystal material having negative anisotropy of dielectric constant (T_{ni}=75° C., Δn=0.10, Δε=-3.2). The liquid crystal material contained 0.6 wt % of an azobenzene group-containing bifunctional

monomer represented by the following formula (2-5). The liquid crystal material injection was followed by heating at 130° C. and quenching. The liquid crystal cell was irradiated with unpolarized ultraviolet light having a wavelength of 365 nm for 30 minutes (irradiation dose: 3 J/cm²) with a cell temperature of 35° C. to polymerize the monomer, and thereby a 4D-RTN mode liquid crystal cell was obtained.



Examples 3-2 to 3-5

[0160] 4D-RTN mode liquid crystal cells of Examples 3-2, 3-3, 3-4, and 3-5 were produced as in Example 3-1, except that the amount m of the initiator functional group-containing diamine unit introduced into the polyamic acid was changed to 20 mol %, 30 mol %, 40 mol %, and 50 mol %, respectively.

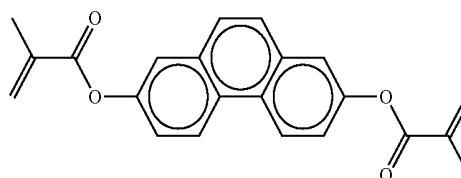
Comparative Example 3-1

[0161] A 4D-RTN mode liquid crystal cell of Comparative Example 3-1 was produced as in Example 3-1, except that the amount m of the initiator functional group-containing diamine unit introduced into the polyamic acid was changed to 0 mol %.

Comparative Example 3-2

[0162] A 4D-RTN mode liquid crystal cell of Comparative Example 3-2 was produced as in Comparative Example 3-1, except that the liquid crystal material contained 0.6 wt % of a bifunctional monomer represented by the formula (M-2) instead of the monomer represented by the formula (2-5).

(M-2)



Comparative Examples 3-3 to 3-7

[0163] 4D-RTN mode liquid crystal cells of Comparative Examples 3-3, 3-4, 3-5, 3-6, and 3-7 were produced as in Examples 3-1, 3-2, 3-3, 3-4, and 3-5, respectively, except that the liquid crystal material contained 0.6 wt % of a bifunctional monomer represented by the formula (M-2) instead of the monomer represented by the formula (2-5).

[0164] The liquid crystal cells were subjected to the same evaluations as in Example 1-1 and the other examples. The results are shown in the following Tables 3-1 and 3-2.

TABLE 3-1

Results of using azobenzene group-containing bifunctional monomer and applying polarized UV light at 35° C.							
	Comparative Example 3-1	Example 3-1	Example 3-2	Example 3-3	Example 3-4	Example 3-5	
Initiator introduced (mol %)	m = 0	m = 10	m = 20	m = 30	m = 40	m = 50	
Alignment state	Initial	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned
Contrast ratio	Initial	5000	5000	5000	5000	4900	4400
	100 h	5000	5000	5000	5000	4900	4200
VHR (%)	Initial	99.2	99.2	99.2	99.2	99.2	99.3
	100 h	94.9	98.3	98.6	99.0	99.0	99.1
rDC voltage (mV)	Initial	-20	-20	-20	-20	-20	-20
	100 h	100	20	20	0	0	0

TABLE 3-2

Results of using phenanthrene group-containing bifunctional monomer and applying polarized UV light at 35° C.							
	Comparative Example 3-2	Comparative Example 3-3	Comparative Example 3-4	Comparative Example 3-5	Comparative Example 3-6	Comparative Example 3-7	
Initiator introduced (mol %)	m = 0	m = 10	m = 20	m = 30	m = 40	m = 50	
Alignment state	Initial	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned
Contrast ratio	Initial	5000	5000	5000	5000	4800	4400
	100 h	4700	5000	5000	5000	4800	3900
VHR (%)	Initial	99.4	99.5	99.5	99.4	99.4	99.5
	100 h	96.3	97.7	97.8	98.4	98.9	99.1
rDC voltage (mV)	Initial	-10	-20	-20	-20	-20	-30
	100 h	70	50	50	40	10	20

[0165] When an azobenzene-containing monomer represented by the formula (2-5) was used, unpolarized ultraviolet light was applied at 35° C. (common PSA conditions) (Table 3-1), and a polymerization initiator functional group was introduced into a side chain of a polyamic acid (partially imidized) constituting the vertical photo-alignment film (m=10 to 50), the backlight exposure power-on test did not cause a VHR decrease or a rDC voltage increase. However, with m=50, the contrast ratio decreased. This is presumably because the introduction of 50 mol % of the polymerization initiator group reduced the amount of the vertical photo-functional group introduced to 50 mol %, decreasing the alignment stability (i.e., stability of the pre-tilt angle).

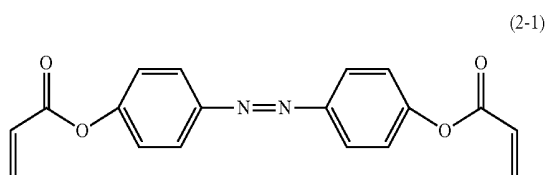
[0166] Table 3-2 shows that when a phenanthrene group-containing bifunctional monomer, which is a conventionally used monomer for PSA, was used (formula (M-2)), the VHR after 100-hour aging decreased as compared with the case of using a monomer represented by the formula (2-5). This is presumably because unreacted monomers of the phenanthrene group-containing bifunctional monomer, having a lower reaction efficiency than the azobenzene group-containing monomer, remained in the liquid crystal layer in a slight amount lower than the detectable level, and were therefore slightly deteriorated under backlight illumination.

Example 4-1

(Production of Liquid Crystal Cell)

[0167] A pair of substrates each having an ITO electrode provided with slits was prepared. A vertical photo-alignment

film was formed on each of the substrates using a polyamic acid containing in a side chain an initiator functional group synthesized as in Condensation Polymerization Example 4 and a vertical functional group. The amount m of the initiator functional group-containing diamine unit introduced into the polyamic acid was 10 mol %. The film formation includes pre-baking the applied alignment agent at 80° C. for five minutes and then post-baking it at 200° C. for 40 minutes. In the present example, the alignment films were not subjected to any special alignment treatment. Still, the vertical alignment can be achieved. A seal was applied to one of the substrates, and beads were scattered on the other substrate. The substrates were then bonded to each other. Between the substrates was injected a liquid crystal material having negative anisotropy of dielectric constant ($T_{ni}=75^{\circ}\text{C.}$, $\Delta n=0.09$, $\Delta\epsilon=-3.0$). The liquid crystal material contained 0.3 wt % of an azobenzene group-containing bifunctional monomer represented by the following formula (2-1). The liquid crystal material injection was followed by heating at 130° C. and quenching. The liquid crystal cell was irradiated with unpolarized ultraviolet light having a wavelength of 365 nm for 30 minutes (irradiation dose: 3 J/cm²) with a cell temperature of 35° C. to polymerize the monomer while a voltage of 5 V was applied to the liquid crystal cell, and thereby a vertical alignment mode liquid crystal cell was obtained.



Examples 4-2 to 4-5

[0168] Vertical alignment mode liquid crystal cells of Examples 4-2, 4-3, 4-4, and 4-5 were produced as in Example 4-1, except that the amount *m* of the initiator functional group-containing diamine unit introduced into the

tained 0.3 wt % of a bifunctional monomer represented by the formula (M-2) instead of the monomer represented by the formula (2-1).

Comparative Examples 4-3 to 4-7

[0171] Vertical alignment mode liquid crystal cells of Comparative Examples 4-3, 4-4, 4-5, 4-6, and 4-7 were produced as in Examples 4-1, 4-2, 4-3, 4-4, and 4-5, respectively, except that the liquid crystal material contained 0.3 wt % of a bifunctional monomer represented by the formula (M-2) instead of the monomer represented by the formula (2-1).

[0172] The liquid crystal cells were subjected to the same evaluations as in Example 1-1 and the other examples. The results are shown in the following Tables 4-1 and 4-2.

TABLE 4-1

Results of using azobenzene group-containing bifunctional monomer and applying polarized UV light at 35° C.							
	Comparative Example 4-1	Example 4-1	Example 4-2	Example 4-3	Example 4-4	Example 4-5	
Initiator introduced (mol %)	<i>m</i> = 0	<i>m</i> = 10	<i>m</i> = 20	<i>m</i> = 30	<i>m</i> = 40	<i>m</i> = 50	
Alignment state	Initial	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned
Contrast ratio	Initial	5000	5000	5000	5000	4900	4900
	100 h	5000	5000	5000	5000	4900	4900
VHR (%)	Initial	99.5	99.5	99.5	99.5	99.5	99.5
	100 h	97.4	99.1	99.1	99.1	99.2	99.1
rDC voltage (mV)	Initial	0	0	-10	-10	-30	-30
	100 h	60	10	0	0	10	0

TABLE 4-2

Results of using phenanthrene group-containing bifunctional monomer and applying polarized UV light at 35° C.							
	Comparative Example 4-2	Comparative Example 4-3	Comparative Example 4-4	Comparative Example 4-5	Comparative Example 4-6	Comparative Example 4-7	
Initiator introduced (mol %)	<i>m</i> = 0	<i>m</i> = 10	<i>m</i> = 20	<i>m</i> = 30	<i>m</i> = 40	<i>m</i> = 50	
Alignment state	Initial	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned	Substantially vertically aligned
Contrast ratio	Initial	5000	5000	5000	5000	5000	5000
	100 h	4800	5000	5000	5000	4900	5000
VHR (%)	Initial	99.3	99.3	99.4	99.4	99.4	99.5
	100 h	98.3	98.8	98.9	99.1	99.1	99.2
rDC voltage (mV)	Initial	-20	-20	-30	-30	-30	-30
	100 h	50	0	0	0	0	-10

polyamic acid was changed to 20 mol %, 30 mol %, 40 mol %, and 50 mol %, respectively.

Comparative Example 4-1

[0169] A vertical alignment mode liquid crystal cell of Comparative Example 4-1 was produced as in Example 4-1, except that the amount *m* of the initiator functional group-containing diamine unit introduced into the polyamic acid was changed to 0 mol %.

Comparative Example 4-2

[0170] A vertical alignment mode liquid crystal cell of Comparative Example 4-2 was produced as in Comparative Example 4-1, except that the liquid crystal material con-

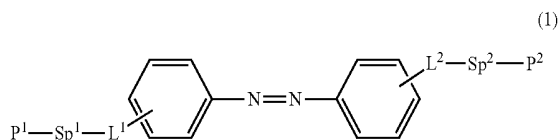
[0173] When an azobenzene-containing monomer represented by the formula (2-1) was used, unpolarized ultraviolet light was applied at 35° C. (with voltage applied) (Table 4-1), and a polymerization initiator functional group was introduced into a side chain of a polyamic acid (partially imidized) constituting the vertical photo-alignment film (*m*=10 to 50), the backlight exposure turn-on test did not cause a VHR decrease or a rDC voltage increase.

[0174] Table 4-2 shows that when a phenanthrene group-containing bifunctional monomer, which is a conventionally used monomer for PSA, was used (formula (M-2)), the VHR after 100-hour aging decreased as compared with the case of using a monomer represented by the formula (2-1). This is presumably because unreacted monomers of the phenanthrene group-containing bifunctional monomer, having a lower reaction efficiency than the azobenzene group-con-

taining monomer, remained in the liquid crystal layer in a slight amount lower than the detectable level, and were therefore slightly deteriorated under backlight illumination.

ADDITIONAL REMARKS

[0175] One aspect of the present invention is directed to a liquid crystal display device including: a pair of substrates; a liquid crystal layer held between the substrates; an alignment film disposed on a liquid crystal layer side surface of at least one of the substrates; and a polymer layer disposed between the liquid crystal layer and the alignment film, the liquid crystal layer containing liquid crystal compounds aligned in a predetermined direction with no voltage applied, the alignment film containing a first polymer containing in its main chain at least one selected from a polyamic acid structure and a polyimide structure, the polymer layer containing a second polymer obtained by polymerizing at least one monomer including at least one monomer represented by the following formula (1), the second polymer covalently bonded to the first polymer,

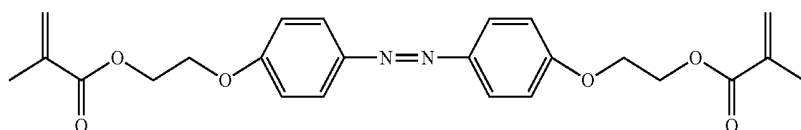
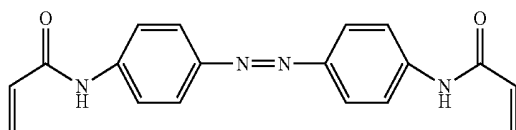
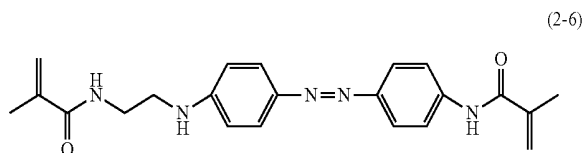
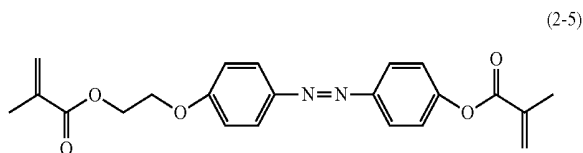
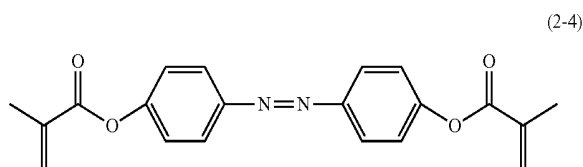
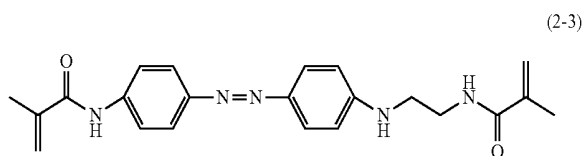
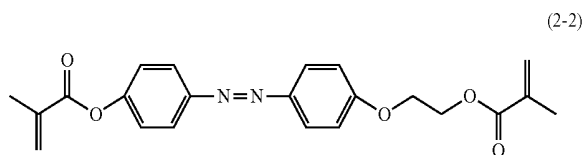
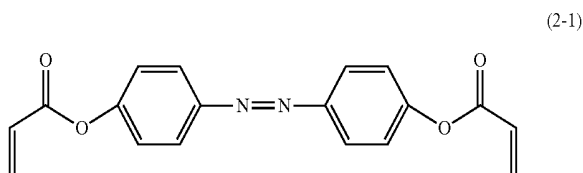


wherein P^1 and P^2 are the same as or different from each other, and each represent an acryloyloxy, methacryloyloxy,

acryloylamino, methacryloylamino, vinyl, or vinyloxy group, Sp^1 and Sp^2 are the same as or different from each other, and each represent a C1-C10 linear, branched, or cyclic alkylene or alkenylene group, or a direct bond, L^1 and L^2 are the same as or different from each other, and each represent a $-\text{NH}-$ group, a $-\text{N}(\text{CH}_3)-$ group, a $-\text{O}-$ group, a $-\text{S}-$ group, or a direct bond, and at least one hydrogen atom in each phenylene group is optionally replaced.

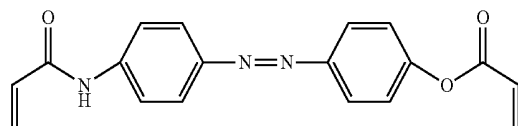
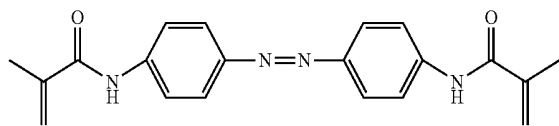
[0176] In the above aspect of the liquid crystal display device, the second polymer in the polymer layer is covalently bonded to the first polymer in the alignment films. This increases the formation rate of the polymer layer (polymerization rate of the monomer), reducing the ultraviolet light irradiation dose for the liquid crystal layer in the polymer layer formation. Also, the polymer layer can be easily formed since it contains the second polymer obtained by polymerizing at least one monomer including at least one monomer represented by the formula (1), further reducing the ultraviolet light irradiation dose for the liquid crystal layer **40** in the polymer layer formation. This is because a polymer of the monomer (1) is likely to separate from the liquid crystal layer **40**. These conditions can prevent a VHR decrease and a residual DC voltage increase even when the liquid crystal display device of the present embodiment is used for a long period of time.

[0177] The at least one monomer represented by the formula (1) may include at least one monomer represented by any of the following formulas (2-1) to (2-17).

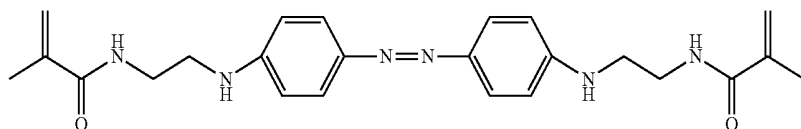


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(2-9)

(2-10)

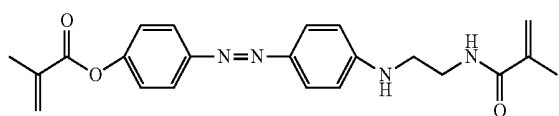


(2-11)



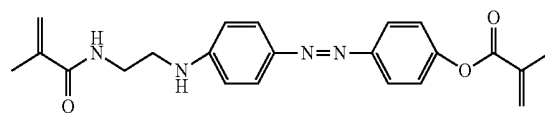
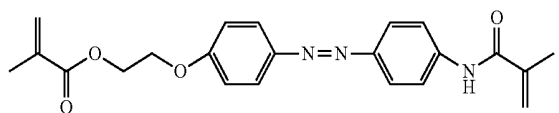
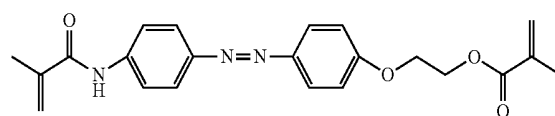
(2-12)

(2-13)

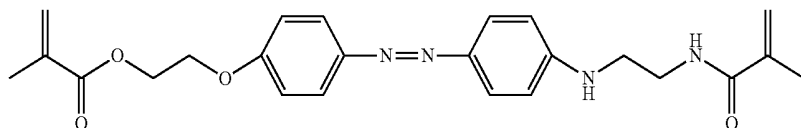


(2-14)

(2-15)



(2-16)

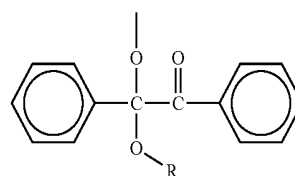


(2-17)

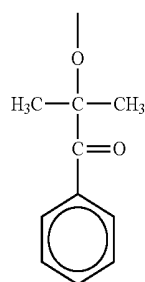
[0178] The first polymer may contain at least one functional group represented by any of the following formulas (3-1) to (3-7):

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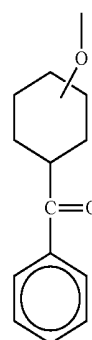
(3-2)



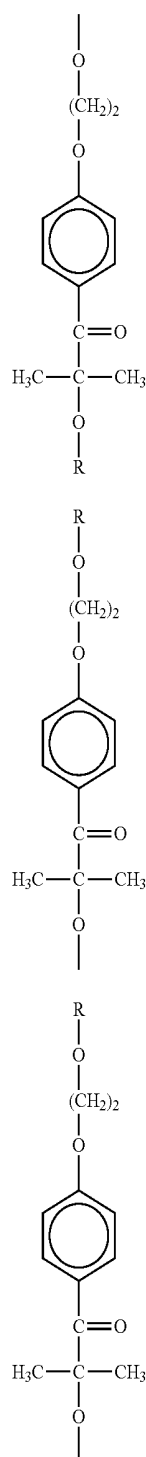
(3-3)



(3-1)

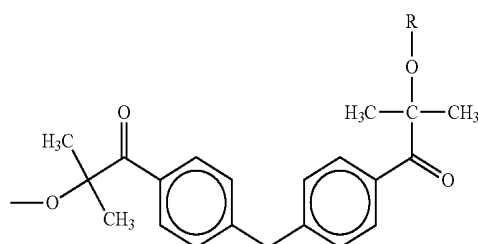


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(3-4)

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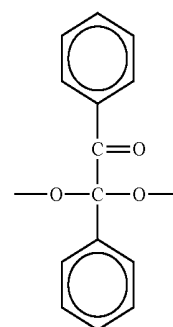
(3-7)

wherein Rs are the same as or different from each other, and each represent a C1-C40 saturated or unsaturated hydrocarbon group.

[0179] The first polymer may contain a functional group represented by the formula (3-1).

[0180] The first polymer may contain a functional group represented by the following formula (4).

(3-5)



(4)

[0181] The first polymer may contain at least one photo-functional group.

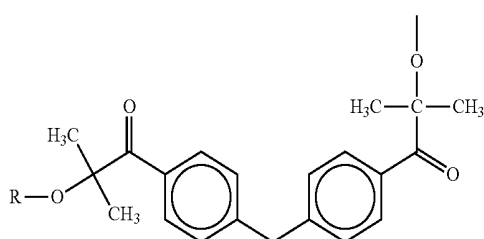
[0182] The first polymer may contain at least one photo-functional group selected from the group consisting of a cinnamate group, an azobenzene group, a chalcone group, and a coumarin group, each of which may contain a substituent.

[0183] The liquid crystal display device may be in a liquid crystal driving mode of a horizontal alignment mode, and the polymer layer may align the liquid crystal compounds in a directional parallel to surfaces of the substrates.

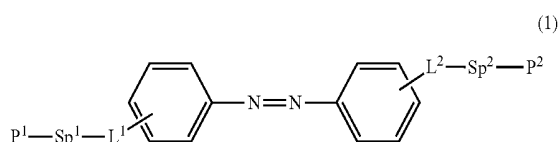
[0184] The liquid crystal display device may be in a liquid crystal driving mode of an FFS mode, an IPS mode, an MVA mode, or a 4D-RTN mode.

[0185] Another aspect of the present invention is directed to a method for manufacturing a liquid crystal display device including a liquid crystal layer containing liquid crystal compounds aligned in a predetermined direction with no voltage applied, the method including: preparing a pair of substrates; forming an alignment film by applying to a surface of at least one of the substrates an alignment agent that contains a first polymer containing at least one selected from a polyamic acid structure and a polyimide structure in its main chain; forming a liquid crystal layer by sealing, between the substrates on at least one of which the alignment film is formed, a liquid crystal composition containing a liquid crystal material and at least one monomer including at least one monomer represented by the following formula (1); and forming a polymer layer from a second polymer

(3-6)



between the alignment film and the liquid crystal layer by irradiating the liquid crystal layer with ultraviolet light, the second polymer being obtained by polymerizing the at least one monomer including at least one monomer represented by the following formula (1), the forming a polymer layer including covalently bonding the second polymer to the first polymer,

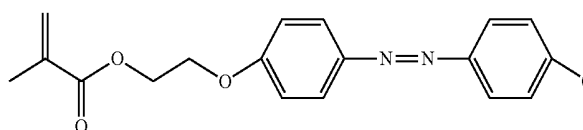
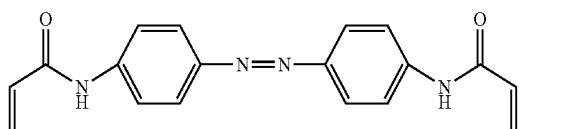
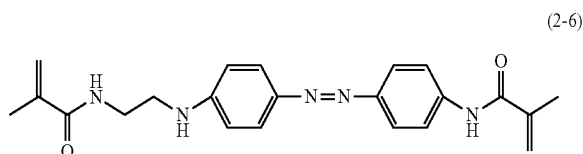
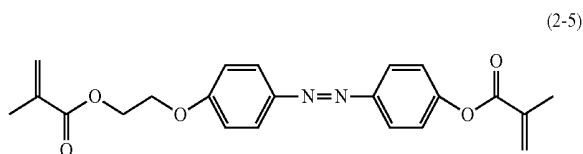
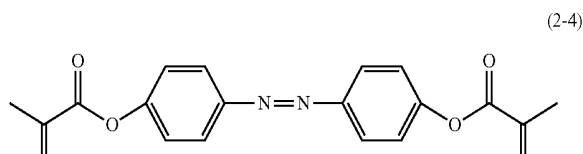
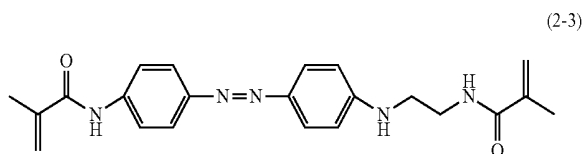
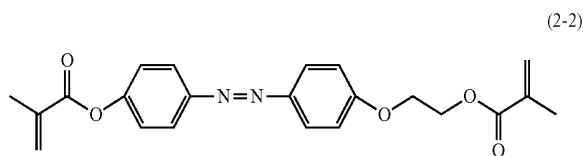
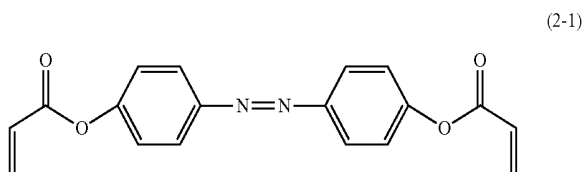


wherein P^1 and P^2 are the same as or different from each other, and each represent an acryloyloxy, methacryloyloxy, acryloylamino, methacryloylamino, vinyl, or vinyloxy group, Sp^1 and Sp^2 are the same as or different from each other, and each represent a C1-C10 linear, branched, or cyclic alkylene or alkenylene group, or a direct bond, L^1 and L^2 are the same as or different from each other, and each

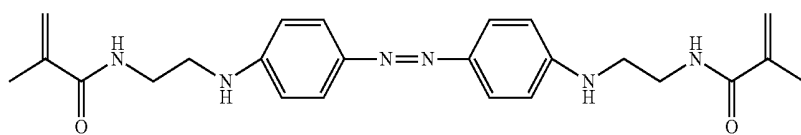
represent a $-\text{NH}-$ group, a $-\text{N}(\text{CH}_3)-$ group, a $-\text{O}-$ group, a $-\text{S}-$ group, or a direct bond, and at least one hydrogen atom in each phenylene group is optionally replaced.

[0186] In the above aspect of the method for manufacturing a liquid crystal display device, the second polymer constituting the polymer layer is covalently bonded to the first polymer constituting the alignment films. This increases the formation rate of the polymer layer (polymerization rate of the monomer), reducing the ultraviolet light irradiation dose for the liquid crystal layer in the polymer layer formation. Also, the polymer layer can be easily formed since it contains the second polymer obtained by polymerizing at least one monomer including at least one monomer represented by the formula (1), further reducing the ultraviolet light irradiation dose for the liquid crystal layer 40 in the polymer layer formation. This is because a polymer of the monomer (1) is likely to separate from the liquid crystal layer 40. These conditions can prevent a VHR decrease and a residual DC voltage increase even when the liquid crystal display device of the present embodiment is used for a long period of time.

[0187] The at least one monomer represented by the formula (1) may include at least one monomer represented by any of the following formulas (2-1) to (2-17).

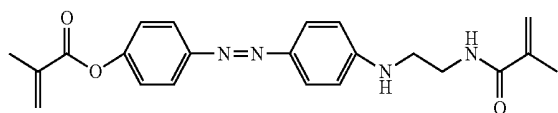


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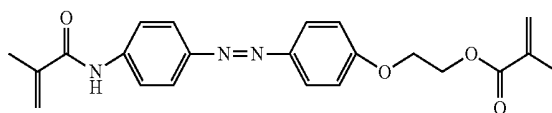


(2-11)

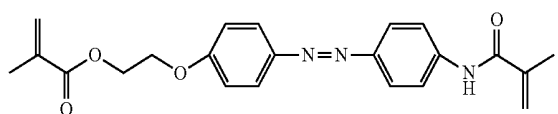
(2-12)



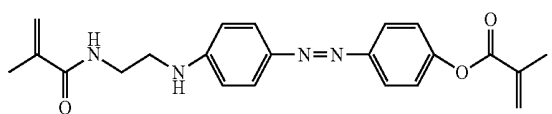
(2-13)



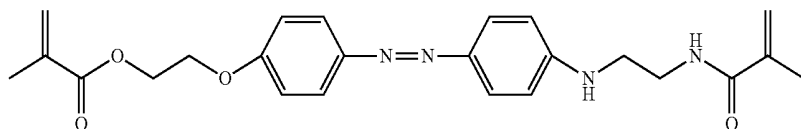
(2-14)



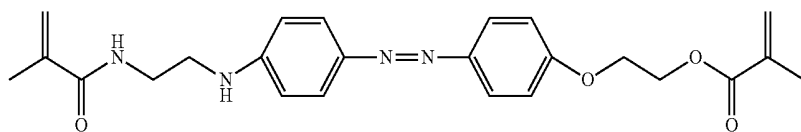
(2-15)



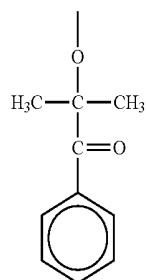
(2-16)



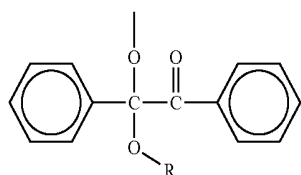
(2-17)



[0188] The first polymer contains at least one functional group represented by any of the following formulas (3-1) to (3-7):

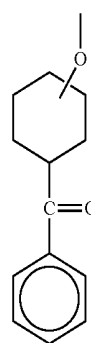


(3-1)

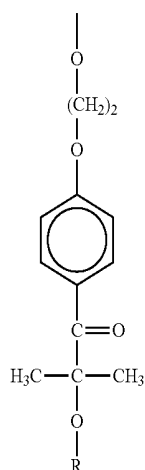


(3-2)

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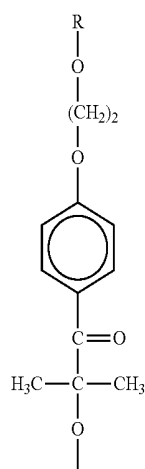


(3-3)

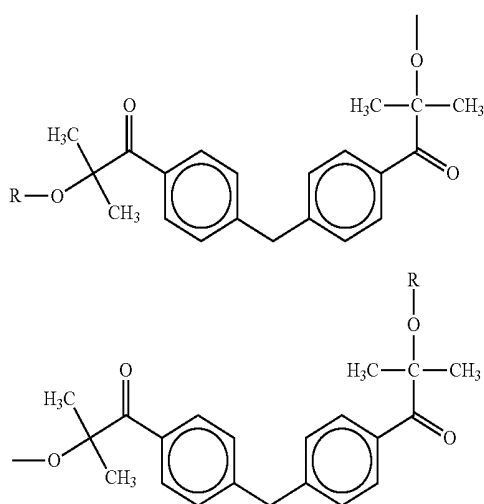


(3-4)

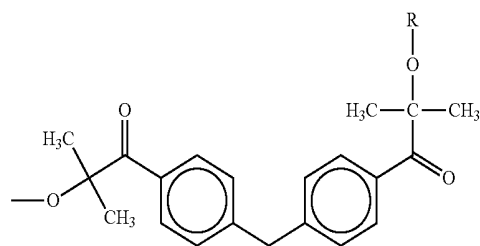
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(3-5)



(3-6)

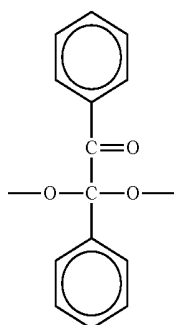


(3-7)

wherein Rs are the same as or different from each other, and each represent a C1-C40 saturated or unsaturated hydrocarbon group.

[0189] The first polymer may contain a functional group represented by the formula (3-1).

[0190] The first polymer may contain a functional group represented by the following formula (4).



(4)

[0191] The first polymer may contain at least one photo-functional group.

[0192] The forming a polymer layer may include irradiating the liquid crystal layer with polarized ultraviolet light so as to polymerize the at least one monomer including at least one monomer represented by the formula (1).

[0193] The forming a polymer layer may include irradiating the liquid crystal layer with unpolarized ultraviolet light so as to polymerize the at least one monomer including at least one monomer represented by the formula (1).

What is claimed is:

1. A liquid crystal display device comprising:

a pair of substrates;

a liquid crystal layer held between the substrates;

an alignment film disposed on a liquid crystal layer side surface of at least one of the substrates; and

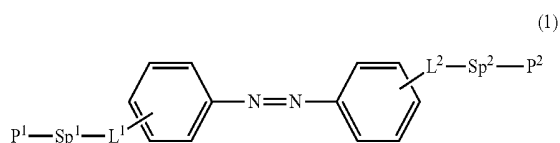
a polymer layer disposed between the liquid crystal layer and the alignment film,

the liquid crystal layer containing liquid crystal compounds aligned in a predetermined direction with no voltage applied,

the alignment film containing a first polymer containing in its main chain at least one selected from a polyamic acid structure and a polyimide structure,

the polymer layer containing a second polymer obtained by polymerizing at least one monomer including at least one monomer represented by the following formula (1),

the second polymer covalently bonded to the first polymer,



(1)

wherein P¹ and P² are the same as or different from each other, and each represent an acryloyloxy, methacryloyloxy, acryloylamino, methacryloylamino, vinyl, or vinyloxy group,

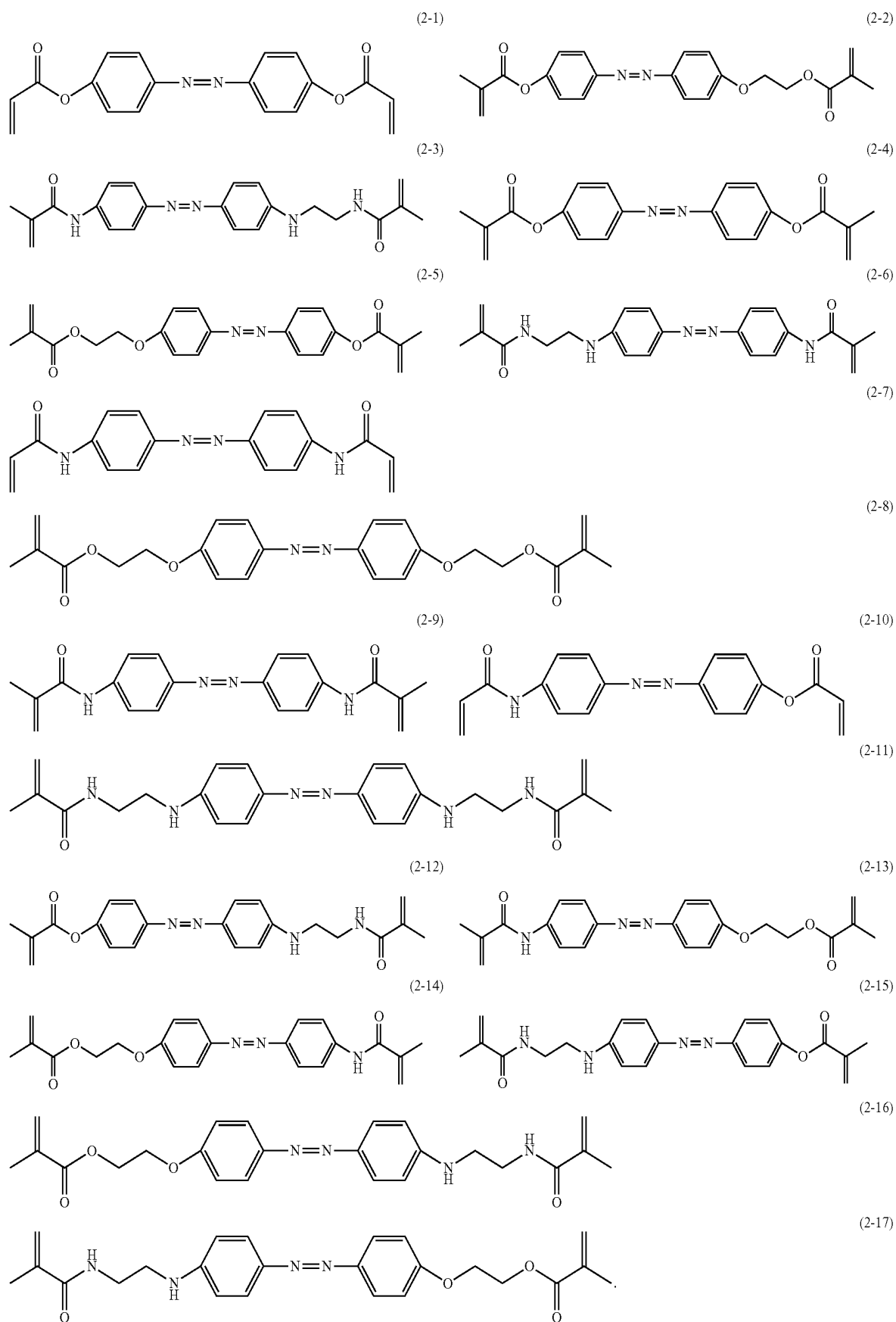
Sp¹ and Sp² are the same as or different from each other, and each represent a C1-C10 linear, branched, or cyclic alkylene or alkenylene group, or a direct bond,

L¹ and L² are the same as or different from each other, and each represent a —NH— group, a —N(CH₃)— group, a —O— group, a —S— group, or a direct bond, and

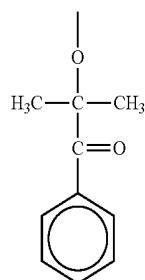
at least one hydrogen atom in each phenylene group is optionally replaced.

2. The liquid crystal display device according to claim 1,

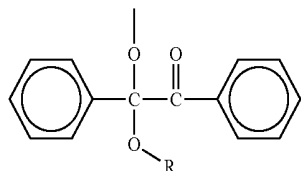
wherein the at least one monomer represented by the formula (1) includes at least one monomer represented by any of the following formulas (2-1) to (2-17):



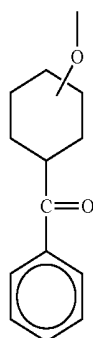
3. The liquid crystal display device according to claim 1, wherein the first polymer contains at least one functional group represented by any of the following formulas (3-1) to (3-7):



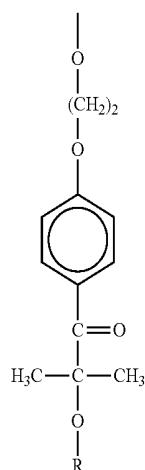
(3-1)



(3-2)

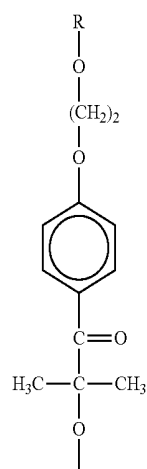


(3-3)

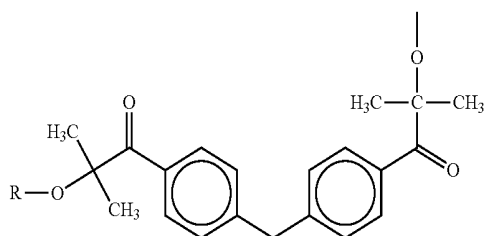


(3-4)

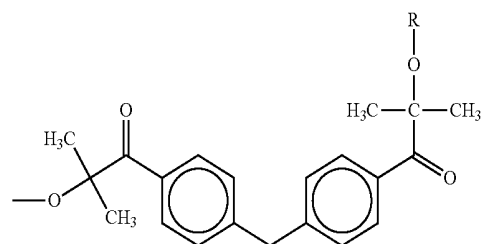
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(3-5)



(3-6)

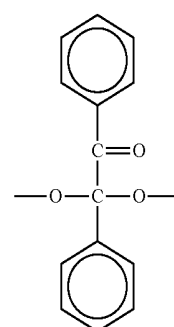


(3-7)

wherein Rs are the same as or different from each other, and each represent a C1-C40 saturated or unsaturated hydrocarbon group.

4. The liquid crystal display device according to claim 3, wherein the first polymer contains a functional group represented by the formula (3-1).

5. The liquid crystal display device according to claim 1, wherein the first polymer contains a functional group represented by the following formula (4):



(4)

6. The liquid crystal display device according to claim 1, wherein the first polymer contains at least one photo-functional group.

7. The liquid crystal display device according to claim 6, wherein the first polymer contains at least one photo-functional group selected from the group consisting of a cinnamate group, an azobenzene group, a chalcone group, and a coumarin group, each of which may contain a substituent.

8. The liquid crystal display device according to claim 1, wherein the liquid crystal display device is in a liquid crystal driving mode of a horizontal alignment mode, and

the polymer layer aligns the liquid crystal compounds in a directional parallel to surfaces of the substrates.

9. The liquid crystal display device according to claim 1, wherein the liquid crystal display device is in a liquid crystal driving mode of an FFS mode, an IPS mode, an MVA mode, or a 4D-RTN mode.

10. A method for manufacturing a liquid crystal display device including a liquid crystal layer containing liquid crystal compounds aligned in a predetermined direction with no voltage applied, the method comprising:

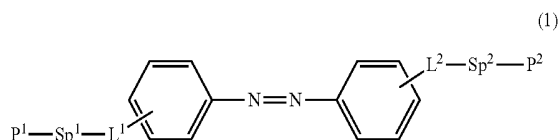
preparing a pair of substrates;

forming an alignment film by applying to a surface of at least one of the substrates an alignment agent that contains a first polymer containing at least one selected from a polyamic acid structure and a polyimide structure in its main chain;

forming a liquid crystal layer by sealing, between the substrates on at least one of which the alignment film is formed, a liquid crystal composition containing a liquid crystal material and at least one monomer including at least one monomer represented by the following formula (1); and

forming a polymer layer from a second polymer between the alignment film and the liquid crystal layer by irradiating the liquid crystal layer with ultraviolet light, the second polymer being obtained by polymerizing the at least one monomer including at least one monomer represented by the following formula (1),

the forming a polymer layer including covalently bonding the second polymer to the first polymer,



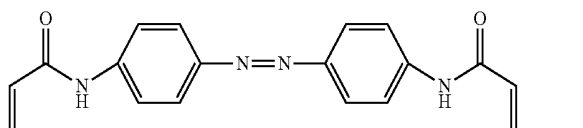
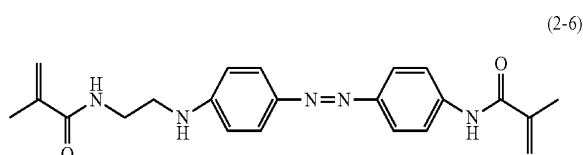
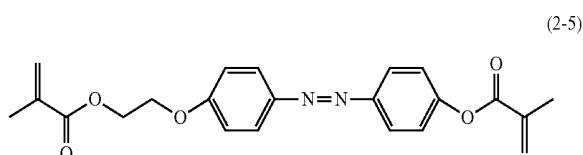
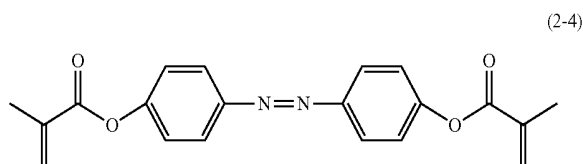
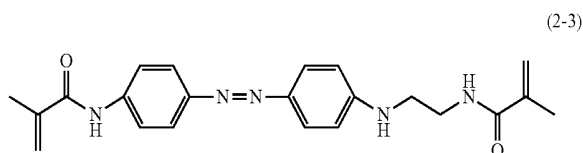
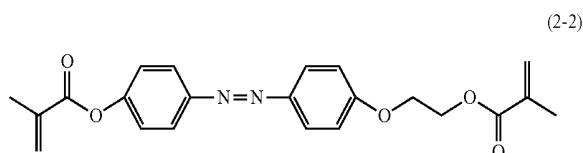
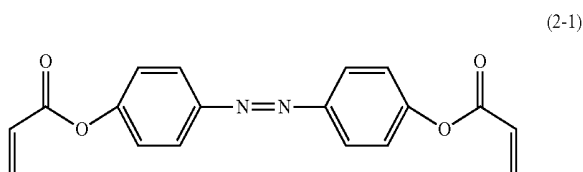
wherein P^1 and P^2 are the same as or different from each other, and each represent an acryloyloxy, methacryloyloxy, acryloylamino, methacryloylamino, vinyl, or vinyloxy group,

Sp^1 and Sp^2 are the same as or different from each other, and each represent a C1-C10 linear, branched, or cyclic alkylene or alkenylene group, or a direct bond,

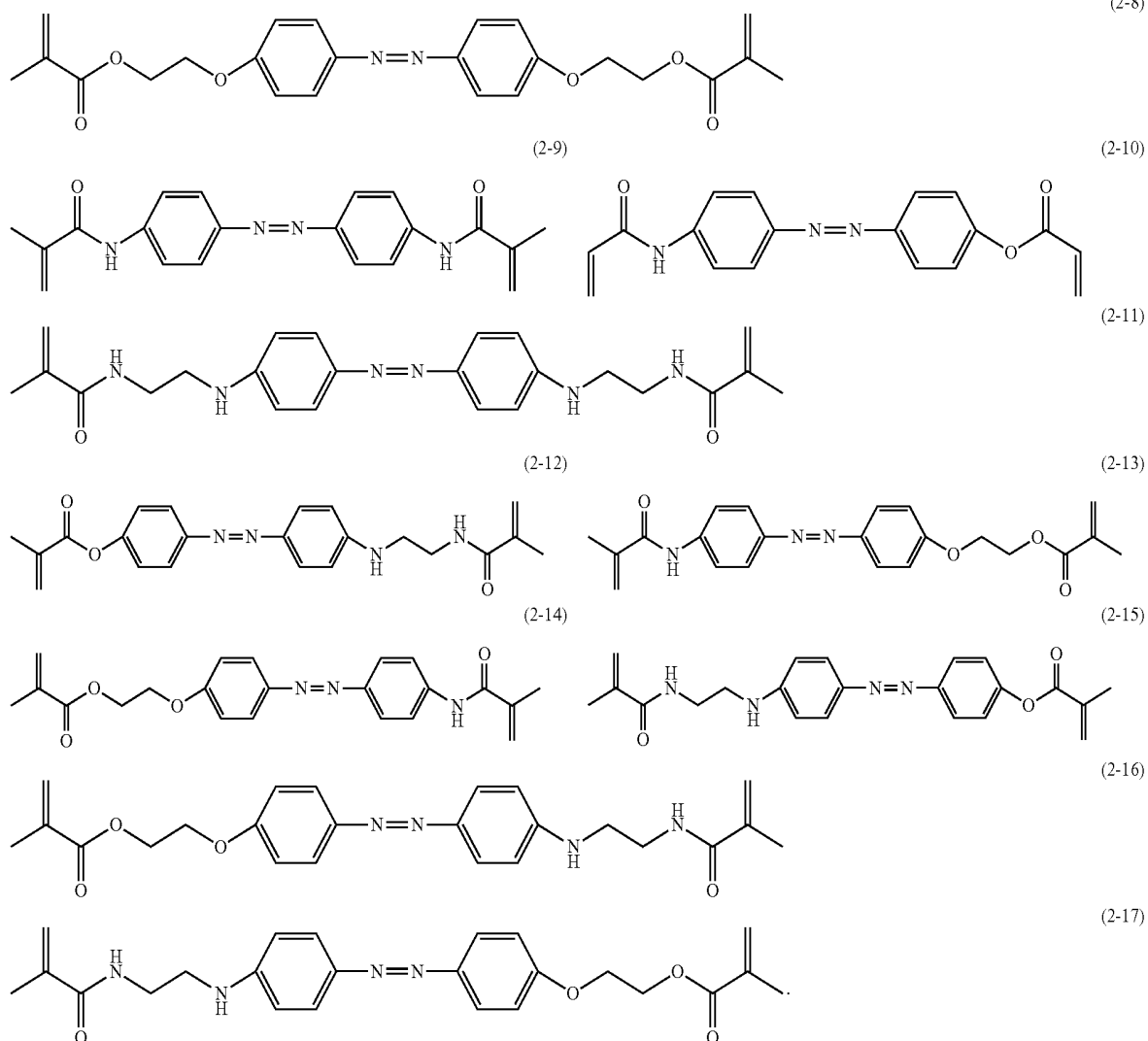
L^1 and L^2 are the same as or different from each other, and each represent a $-\text{NH}-$ group, a $-\text{N}(\text{CH}_3)-$ group, a $-\text{O}-$ group, a $-\text{S}-$ group, or a direct bond, and at least one hydrogen atom in each phenylene group is optionally replaced.

11. The method for manufacturing a liquid crystal display device according to claim 10,

wherein the at least one monomer represented by the formula (1) includes at least one monomer represented by any of the following formulas (2-1) to (2-17):

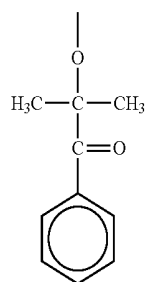


-continued

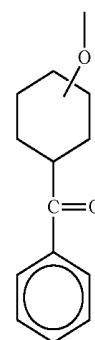
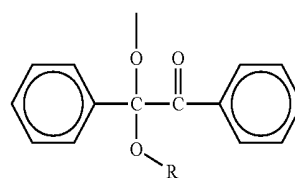


12. The method for manufacturing a liquid crystal display device according to claim 10,

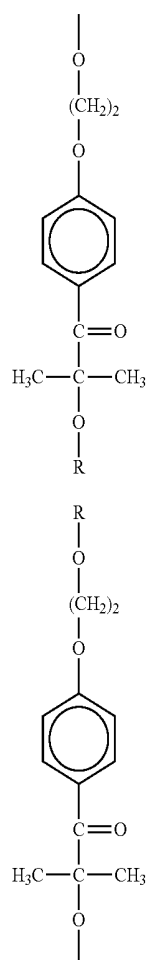
wherein the first polymer contains at least one functional group represented by any of the following formulas (3-1) to (3-7):



-continued



-continued



(3-4)

wherein Rs are the same as or different from each other, and each represent a C1-C40 saturated or unsaturated hydrocarbon group.

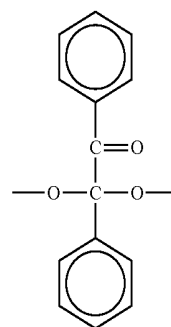
13. The method for manufacturing a liquid crystal display device according to claim **12**,

wherein the first polymer contains a functional group represented by the formula (3-1).

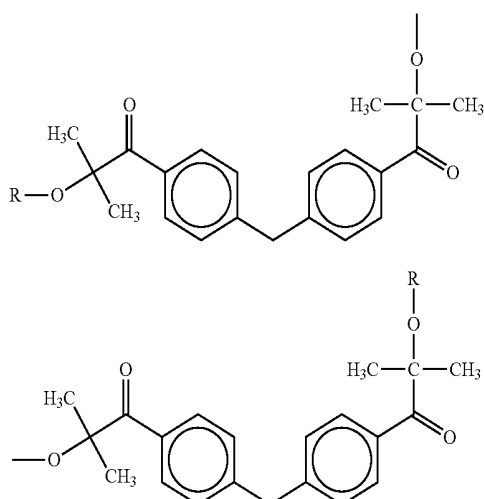
14. The method for manufacturing a liquid crystal display device according to claim **10**,

wherein the first polymer contains a functional group represented by the following formula (4):

(3-5)



(4)



(3-6)

15. The method for manufacturing a liquid crystal display device according to claim **10**,

wherein the first polymer contains at least one photo-functional group.

16. The method for manufacturing a liquid crystal display device according to claim **10**,

wherein the forming a polymer layer includes irradiating the liquid crystal layer with polarized ultraviolet light so as to polymerize the at least one monomer including at least one monomer represented by the formula (1).

17. The method for manufacturing a liquid crystal display device according to claim **10**,

wherein the forming a polymer layer includes irradiating the liquid crystal layer with unpolarized ultraviolet light so as to polymerize the at least one monomer including at least one monomer represented by the formula (1).

* * * * *

专利名称(译)	液晶显示装置及其制造方法		
公开(公告)号	US20200026129A1	公开(公告)日	2020-01-23
申请号	US16/503267	申请日	2019-07-03
[标]申请(专利权)人(译)	夏普株式会社		
申请(专利权)人(译)	夏普株式会社		
当前申请(专利权)人(译)	夏普株式会社		
[标]发明人	MIZUSAKI MASANOBU		
发明人	MIZUSAKI, MASANOBU		
IPC分类号	G02F1/1337 G02F1/1343 C08G73/10 C08L79/08		
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优先权	62/702159 2018-07-23 US		
外部链接	Espacenet USPTO		

摘要(译)

本发明提供一种液晶显示装置，其包括：一对基板；以及一对基板。保持在基板之间的液晶层；取向膜，其配置在至少一个基板的液晶层侧面上。聚合物层和设置在液晶层与取向膜之间的聚合物层，该液晶层包含在没有施加电压的情况下在预定方向上取向的液晶化合物，该取向膜在其主链中包含选自以下的至少一种的第一聚合物 聚合物层包含聚酰胺酸结构和聚酰亚胺结构，该聚合物层包含通过使至少一种包含偶氮苯基的单体聚合而获得的第二聚合物，该第二聚合物与该第一聚合物共价键合。

